

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:02:27 ; Search time 51.308 Seconds
(without alignments)
699.375 Million cell updates/sec

Title: US-10-661-784-3
Perfect score: 637
Sequence: 1 GSKDFVQPTKICVGRD.....VPWEKIIYTVVNHWECEP 127

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A Geneseq_29Jan04:*
1: Geneseqp1980s:*
2: Geneseqp1990s:*
3: Geneseqp2000s:*
4: Geneseqp2001s:*
5: Geneseqp2002s:*
6: Geneseqp2003as:*
7: Geneseqp2003bs:*
8: Geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB ID	Description
1	618	90.0	123	3	AA95426 Human hig
2	618	90.0	304	6	ABP70801 Human ext
3	618	90.0	322	6	ABP70799 Human ext
4	618	90.0	329	6	ABU92044 Human pro
5	618	90.0	358	6	ABP70800 Human ext
6	618	90.0	390	6	ABU99149 Novel hum
7	618	90.0	398	6	ABU99143 Novel hum
8	618	90.0	427	8	ABE76864 Human pro
9	618	90.0	615	6	ABU99144 Novel hum
10	618	90.0	626	5	ABP78707 Human hig
11	618	90.0	644	4	ABG21101 Novel hum
12	618	90.0	644	5	ABP78710 Human hig
13	618	90.0	644	6	ABU99150 Novel hum
14	618	90.0	644	6	ABU99145 Novel hum
15	596	85.3	122	3	ABP37447 Human kin
16	585	85.2	435	4	ABG21105 Novel hum
17	556.5	81.0	117	2	AA333350 Domaine 3
18	440	64.0	436	1	AA40257 Bradykini
19	413	60.1	434	1	AA40633 Bradykini
20	411	59.8	357	6	ABR41302 Human DIT
21	388	56.5	235	5	ABG60077 Human DIT
22	320.5	46.7	248	4	ABG21102 Novel hum
23	316	46.0	369	4	ABG21099 Novel hum
24	190	27.7	305	4	ABG21100 Novel hum
25	171.5	25.0	167	2	AAW98907 Mouse IMC

26	166	24.2	32	3	AA95418 Anti-angi
27	163.5	23.8	126	3	AB37445 Human cys
28	163.5	23.8	145	2	AAW32323 Mature hu
29	163.5	23.8	145	2	AAW31902 Human cys
30	163.5	23.8	145	2	AAW31902 Human cys
31	163.5	23.8	145	4	AAE02410 Human cys
32	163.5	23.8	145	4	AAE04439 Human cys
33	163.5	23.8	145	7	ADD14098 Human src
34	163.5	23.8	167	2	AAV02287 Secreted
35	163	23.7	167	2	ADA45154 Human pol
36	163	23.7	178	7	AAW69734 Human cys
37	161	23.4	32	3	AA95408 Anti-angi
38	157	22.9	122	3	AAW37446 Human kin
39	154.5	22.3	167	2	AAW98910 Mouse IMC
40	153	22.3	27	3	AA95425 Anti-angi
41	144.5	21.0	121	3	AA81200 Human mut
42	144.5	21.0	128	3	AA81189 Human mut
43	143.5	20.9	121	3	AA81198 Human mut
44	143.5	20.9	128	3	AA81187 Human mut
45	142.5	20.7	118	3	AA81218 Bovine mu

ALIGNMENTS

RESULT 1
AA95426
ID AA95426 standard; peptide; 123 AA.
XX AC AA95426;
XX DT 25-SEP-2000 (first entry)
XX DE Human high mol.wt. kininogen domain 3.
XX KW Anti-angiogenic; angiogenesis; inhibitor; kininogen; homologue;
XX KW endothelial cell proliferation; apoptosis; cancer; ocular disorder;
XX KW rheumatoid arthritis; cytostatic; antiarthritic; antirheumatic; therapy;
XX KW human; D3 peptide.
XX OS Homo sapiens.
XX PN WO200035407-A2.
XX PD 22-JUN-2000.
XX PF 02-DEC-1999; 99WO-US028465.
XX PR 16-DEC-1998; 98US-0112427P.
XX PA (UTEM) UNIV TEMPLE.
XX PA (MCCR/) MCCRAE R K.
XX PI McCrae RK;
XX DR WPI; 2000-442247/39.
XX PT Composition for inhibiting angiogenesis and endothelial cell
XX PT proliferation, inducing endothelial cell apoptosis and creating cancer, 3
XX PT rheumatoid arthritis, and ocular disorders comprises a kininogen domain 3
XX PT analog.
XX PS Disclosure; Page 4; 44pp; English.
XX CC The present sequence is that of domain 3 of human high mol.wt. kininogen
XX CC (HK). The invention provides peptides (see AA95405-24) that are
XX CC analogues of certain sites in the HK domain 3, specifically Asn275-
XX CC Lys282, Cys246-Cys249, Leu331-Tyr338 and Tyr299-Ser314. The peptides, in
XX CC which native Cys residues may be replaced by Ala residues, inhibit
XX CC endothelial cell proliferation and may also induce endothelial cell
XX CC apoptosis. Compositions including the peptides are used in claimed
XX CC methods for inhibiting angiogenesis, inhibiting endothelial cell
XX CC proliferation, and inducing endothelial cell apoptosis. Cancer,

CC rheumatoid arthritis, and ocular disorders characterized by undesired
 CC vascularization of the retina are treated
 XX
 SQ Sequence 123 AA;
 Query Match 90.0%; Score 618; DB 3; Length 123;
 Best Local Similarity 100.0%; Pred. No. 5.5e-63;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDIPKSPLEELTHITIKLNAENNAATFYFKIDNVKKARVQV 62
 DB 1 GKDFVQPTKICVGCPRDIPKSPLEELTHITIKLNAENNAATFYFKIDNVKKARVQV 60
 QY 63 AGKKYFIDFVARETTCKESNEELTESCTKLGQSLDCNAEVVYVPEKKIYPTV 118
 DB 61 AGKKYFIDFVARETTCKESNEELTESCTKLGQSLDCNAEVVYVPEKKIYPTV 116
 RESULT 2
 ABP70801
 ID ABP70801 standard; protein; 304 AA.
 XX
 AC ABP70801;
 DT 26-AUG-2003 (first entry)
 XX
 DE Human extracellular messenger, EXMES-28.
 XX
 KW Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO2003018612-A2.
 XX
 PD 06-MAR-2003.
 XX
 PF 22-AUG-2002; 2002WO-US027213.
 XX
 PR 24-AUG-2001; 2001US-0314811P.
 PR 14-DEC-2001; 2001US-0340584P.
 PR 18-JAN-2002; 2002US-0350595P.
 PR 11-MAR-2002; 2002US-0363432P.
 PR 15-MAR-2002; 2002US-0364607P.
 PR 05-APR-2002; 2002US-0370761P.
 PR 24-JUN-2002; 2002US-0391378P.
 XX
 PA (INCY-) INCYTE GENOMICS INC.
 XX
 PI Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebarradian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Ramkumar J;
 XX
 DR WPI; 2003-278643/27.
 DR N-PSDB; ACC42386.
 XX
 PT New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 PT disorders or cancer.
 XX
 PS Claim 1; Page 207; 224pp; English.
 XX
 CC The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to-28; ABP70774-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
 CC for diagnosing or treating a disease or condition associated with
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer

XX
 SQ Sequence 304 AA;
 Query Match 90.0%; Score 618; DB 6; Length 304;
 Best Local Similarity 100.0%; Pred. No. 1.9e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDIPKSPLEELTHITIKLNAENNAATFYFKIDNVKKARVQV 62
 DB 130 GKDFVQPTKICVGCPRDIPKSPLEELTHITIKLNAENNAATFYFKIDNVKKARVQV 189
 QY 63 AGKKYFIDFVARETTCKESNEELTESCTKLGQSLDCNAEVVYVPEKKIYPTV 118
 DB 190 AGKKYFIDFVARETTCKESNEELTESCTKLGQSLDCNAEVVYVPEKKIYPTV 245
 RESULT 3
 ABP70799
 ID ABP70799 standard; protein; 322 AA.
 XX
 AC ABP70799;
 DT 26-AUG-2003 (first entry)
 XX
 DE Human extracellular messenger, EXMES-26.
 XX
 KW Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO2003018612-A2.
 XX
 PD 06-MAR-2003.
 XX
 PF 22-AUG-2002; 2002WO-US027213.
 XX
 PR 24-AUG-2001; 2001US-0314811P.
 PR 14-DEC-2001; 2001US-0340584P.
 PR 18-JAN-2002; 2002US-0350595P.
 PR 11-MAR-2002; 2002US-0363432P.
 PR 15-MAR-2002; 2002US-0364607P.
 PR 05-APR-2002; 2002US-0370761P.
 PR 24-JUN-2002; 2002US-0391378P.
 XX
 PA (INCY-) INCYTE GENOMICS INC.
 XX
 PI Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebarradian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Ramkumar J;
 XX
 DR WPI; 2003-278643/27.
 DR N-PSDB; ACC42386.
 XX
 PT New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 PT disorders or cancer.
 XX
 PS Claim 1; Page 205-206; 224pp; English.
 XX
 CC The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to-28; ABP70774-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
 CC for diagnosing or treating a disease or condition associated with
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer

CC	polynucleotides. The diseases or conditions associated with decreased expression or overexpression of PMM are cell proliferation disorders (e.g. cancer, atherosclerosis), neurological disorders (e.g. epilepsy, Huntington's disease, stroke), immune/inflammatory disorders (e.g. AIDS, allergies), developmental disorders (e.g. hypothyroidism, Cushing's syndrome), gastrointestinal or epithelial disorders, and infections. The PMM polypeptides or their fragments are useful in screening compounds for effectiveness as agonists or antagonists of the polypeptides, or in altering the expression of the target polynucleotide and compounds that specifically bind to, or modulate the activity of the polypeptide.
CC	ABU92041-ABU92060 represent the human PMM polypeptides of the invention
XX	
SQ	Sequence 329 AA;
	Query Match 90.0%; Score 618; DB 6; Length 329;
	Best Local Similarity 100.0%; Pred. No. 2.1e-62;
	Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	3 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITKLNAENNAIFYKIDNVKKARQVQV 62
DB	155 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITKLNAENNAIFYKIDNVKKARQVQV 214
QY	63 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNNAEYVVPWEKKIYPTV 118
DB	215 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNNAEYVVPWEKKIYPTV 270
RESULT 5	
ABP70800	
ID	ABP70800 standard; protein; 358 AA.
XX	
AC	ABP70800;
XX	
DT	26-AUG-2003 (first entry)
XX	
DE	Human extracellular messenger, EXMES-27.
XX	
KW	Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
KW	immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
KW	endocrine disorder; cancer.
XX	
OS	Homo sapiens.
XX	
XX	WO2003018612-A2.
PN	
PD	06-MAR-2003.
XX	
PF	22-AUG-2002; 2002WO-US027213.
XX	
PR	24-AUG-2001; 2001US-0314811P.
PR	14-DEC-2001; 2001US-0340584P.
PR	18-JAN-2002; 2002US-0350595P.
PR	11-MAR-2002; 2002US-0363432P.
PR	15-MAR-2002; 2002US-0364607P.
PR	05-APR-2002; 2002US-0370761P.
PR	24-JUN-2002; 2002US-0391378P.
XX	
XX	(INCY-) INCYTE GENOMICS INC.
PA	
XX	Duggan BM, Lee S, Baughn MR, Hafalia AJA, Walia NK, Elliott VS;
PI	Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang Y, Burford N;
PI	Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebartadian Y;
PI	Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
PI	Ramkumar J;
XX	
XX	WPI; 2003-278643/27.
DR	N-PSDB; ACC42387.
XX	
XX	New human extracellular messenger (EXMES) polypeptide, useful for
PT	preparing a composition for treating a disease associated with decreased
PT	expression or overexpression of functional EXMES e.g. autoimmune
PT	disorders or cancer.
XX	

QY	3 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITKLNAENNAIFYKIDNVKKARQVQV 62
DB	148 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITKLNAENNAIFYKIDNVKKARQVQV 207
QY	63 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNNAEYVVPWEKKIYPTV 118
DB	208 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNNAEYVVPWEKKIYPTV 263
RESULT 4	
ABU92044	
ID	ABU92044 standard; protein; 329 AA.
XX	
AC	ABU92044;
XX	
DT	15-JUL-2003 (first entry)
XX	
DE	Human protein modification and maintenance molecule-24 (PMM-24).
XX	
KW	Human; protein modification and maintenance molecule; PMM; cancer;
KW	cell proliferation disorder; atherosclerosis; neurological disorder;
KW	epilepsy; Huntington's disease; stroke; immune disorder; allergy;
KW	inflammatory disorder; AIDS; developmental disorder; hypothyroidism;
KW	Cushing's syndrome; gastrointestinal disorder; epithelial disorder;
KW	infection; cytostatic; antiatherosclerotic; anticonvulsant; nootropic;
KW	neuroprotective; cerebroprotective; anti-HIV; anti-allergic; vulnary;
KW	antiinflammatory; thyromimetic.
XX	
OS	Homo sapiens.
XX	
XX	WO2003031939-A2.
PN	
PD	17-APR-2003.
XX	
PF	11-OCT-2002; 2002WO-US032850.
XX	
PR	12-OCT-2001; 2001US-0329689P.
PR	25-OCT-2001; 2001US-0335703P.
PR	09-NOV-2001; 2001US-0348887P.
PR	28-NOV-2001; 2001US-0334145P.
PR	06-DEC-2001; 2001US-0337451P.
PR	14-DEC-2001; 2001US-0340584P.
XX	
XX	(INCY-) INCYTE GENOMICS INC.
XX	
PI	Ramkumar J, Gorvay AE, Baughn MR, Emerling BM, Yang J, Lee SY;
PI	Tran UK, Becha SD, Duggan BM, Lee EA, Griffin JA, Li JX;
PI	Sprague WM, Hafalia AJA, Chawla NK, Lehr-Mason PM, Kable AE, Yue H;
PI	Marquis JP, Yao MG, Richardson TW, Tang TY, Jin P, Chien D;
PI	Bhatia U, Burrill JD, Lee S, Blake JU, Ho A, Zheng W;
XX	
XX	WPI; 2003-430274/40.
DR	N-PSDB; ACA92439.
XX	
XX	New human protein modification and maintenance molecules (PMM), useful
PT	for diagnosing, treating and preventing diseases or conditions associated
PT	with the aberrant PMM expression e.g. cancer, atherosclerosis, or
PT	infections.
XX	
PS	Claim 1; Page 264-265; 31pp; English.
XX	
CC	The present invention relates to the isolation of human protein
CC	modification and maintenance molecules (PMM), and the polynucleotide
CC	sequences encoding them. A total of 40 PMM polypeptides (designated PMM
CC	-1 to PMM-40) are disclosed. The sequences of the invention are useful
CC	for diagnosing a condition or disease associated with the expression of
CC	PMM in a subject, preparing a polyclonal or monoclonal antibody, and
CC	generating an expression profile of a sample containing the

```

PS  Claim 1; Page 206; 224pp; English.
XX
CC  The present invention relates to novel human extracellular messenger
CC  proteins (EXMES-1 to-28; AB70774-AB70801) and their coding sequences
CC  (ACC4361-ACC4388). The proteins are useful for preparing a composition
CC  for diagnosing or treating a disease or condition associated with
CC  decreased expression or overexpression of functional EXMES e.g.
CC  autoimmune/inflammatory disorders, diabetes, endocrine disorders or
CC  cancer
XX
SQ  Sequence 358 AA;
Query Match          90.0%; Score 618; DB 6; Length 358;
Best Local Similarity 100.0%; Pred. No. 2.4e-62;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY  3 GKDFVQPPKICVCGPRDIPNTSPLEETLTHITIKLAENNAFFYKIDNVKKARQV 62
Db  184 GKDFVQPPKICVCGPRDIPNTSPLEETLTHITIKLAENNAFFYKIDNVKKARQV 243
QY  63 AGKXYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKIIPTV 118
Db  244 AGKXYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKIIPTV 299
RESULT 6
ABU99149
ID  ABU99149 standard; protein; 390 AA.
AC  ABU99149;
XX
DT  01-AUG-2003 (first entry)
DE  Novel human GPCR related protein NOV12g.
KW  Human; G-protein coupled receptor related protein; GPCR related protein;
KW  NOV; cytosolic; cardiant; antiarteriosclerotic; antidiabetic;
KW  immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
KW  antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
KW  NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
KW  diabetes; immune disorder; AIDS; obesity; asthma;
KW  haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
KW  infection; multiple sclerosis; cancer-associated cachexia;
KW  wasting disorder; chronic disease; neurogenesis; cell differentiation;
KW  cell proliferation; haematopoiesis; wound healing; angiogenesis;
KW  chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
XX
OS  Homo sapiens.
XX
PN  WO200299116-A2.
XX
PD  12-DEC-2002.
XX
PF  04-JUN-2001; 2001US-02956072.
PR  04-JUN-2001; 2001US-02956072.
PR  06-JUN-2001; 2001US-0295661P.
PR  06-JUN-2001; 2001US-0295640P.
PR  06-JUN-2001; 2001US-0296418P.
PR  14-JUN-2001; 2001US-0296418P.
PR  15-JUN-2001; 2001US-0298285P.
PR  21-JUN-2001; 2001US-0298556P.
PR  26-JUN-2001; 2001US-0299949P.
PR  28-JUN-2001; 2001US-0300883P.
PR  13-AUG-2001; 2001US-0301550P.
PR  27-AUG-2001; 2001US-0311972P.
PR  29-AUG-2001; 2001US-0315071P.
PR  29-AUG-2001; 2001US-0315660P.
PR  14-SEP-2001; 2001US-0322293P.
PR  17-SEP-2001; 2001US-0322706P.
PR  14-DEC-2001; 2001US-0341186P.
PR  28-FEB-2002; 2002US-0361189P.
PR  12-MAR-2002; 2002US-0363673P.
PR  12-MAR-2002; 2002US-0363676P.
PR
XX
PR  03-JUN-2002; 2002US-00363676.
XX
FA  (CURA-) CURAGEN CORP.
XX
PI  Anderson DW, Baumgartner JC, Boldog FL, Casman SU, Edinger SR;
PI  Gangoli EA, Gerlach VL, Gorman L, Guo X, Hjal T, Kekuda R, Li L;
PI  Macdougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
PI  Pena CE, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet CAM;
PI  Voss EZ, Zerhusen BD;
XX
DR  WPI; 2003-140627/13.
DR  N-PSDB; ACD03653.
XX
XX  New NOVX polypeptides and nucleic acids, useful for preventing or
XX  treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
XX  atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
XX  pharmacogenomics.
XX
PS  Claim 1; Page 147; 332pp; English.
XX
CC  The invention describes an isolated polypeptide (I) comprising any of 27
CC  118-961 residue amino acid sequences, given in the specification, a
CC  mature form of them, a sequence that is at least 95 % identical to them,
CC  or a sequence having one or more conservative substitutions in them. The
CC  polypeptide is useful in manufacturing a medicament for treating a
CC  syndrome associated with a human disease selected from a pathology
CC  associated with the polypeptide. The NOVX polypeptides, polynucleotides
CC  and antibodies are useful in treating or preventing NOVX-associated
CC  disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
CC  disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
CC  disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
CC  associated cachexia, and other wasting disorders associated with chronic
CC  diseases. The nucleic acids and polypeptides may also be used as targets
CC  for the identification of small molecules that modulate or inhibit e.g.
CC  neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
CC  wound healing and angiogenesis, in gene therapy, in generation of
CC  antibodies that bind immunospecifically to NOVX substances for use in
CC  therapeutic or diagnostic methods. The nucleic acids are further used as
CC  hybridisation probes, in chromosome mapping, tissue typing, preventive
CC  medicine, and pharmacogenomics. The polypeptides are also useful as
CC  vaccines. This is the amino acid sequence of a novel human G-protein
CC  coupled receptor related protein NOV
XX
SQ  Sequence 390 AA;
Query Match          90.0%; Score 618; DB 6; Length 390;
Best Local Similarity 100.0%; Pred. No. 2.7e-62;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY  3 GKDFVQPPKICVCGPRDIPNTSPLEETLTHITIKLAENNAFFYKIDNVKKARQV 62
Db  216 GKDFVQPPKICVCGPRDIPNTSPLEETLTHITIKLAENNAFFYKIDNVKKARQV 275
QY  63 AGKXYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKIIPTV 118
Db  276 AGKXYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKIIPTV 331
RESULT 7
ABU99143
ID  ABU99143 standard; protein; 398 AA.
XX
AC  ABU99143;
XX
DT  01-AUG-2003 (first entry)
XX
DE  Novel human GPCR related protein NOV12a.
XX
KW  Human; G-protein coupled receptor related protein; GPCR related protein;
KW  NOV; cytosolic; cardiant; antiarteriosclerotic; antidiabetic;
KW  immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
KW  antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
KW  NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;

```


KW diabetes; immune disorder; AIDS; obesity; asthma;
KW hematopoietic disorder; Parkinson's disease; Alzheimer's disease;
KW infection; multiple sclerosis; cancer-associated cachexia;
KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
KW cell proliferation; hematopoiesis; wound healing; angiogenesis;
KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
XX Homo sapiens.
XX
XX WO200299116-A2.
XX
XX 12-DEC-2002.
XX
XX 04-JUN-2002; 2002WO-US017428.
XX
XX 04-JUN-2001; 2001US-0295607P.
XX 04-JUN-2001; 2001US-029561P.
XX 06-JUN-2001; 2001US-0296404P.
XX 06-JUN-2001; 2001US-0296418P.
XX 14-JUN-2001; 2001US-0298285P.
XX 15-JUN-2001; 2001US-0298556P.
XX 21-JUN-2001; 2001US-0299949P.
XX 26-JUN-2001; 2001US-0300883P.
XX 28-JUN-2001; 2001US-0301550P.
XX 13-AUG-2001; 2001US-0311972P.
XX 27-AUG-2001; 2001US-0315071P.
XX 29-AUG-2001; 2001US-0315660P.
XX 14-SEP-2001; 2001US-0322293P.
XX 17-SEP-2001; 2001US-0322706P.
XX 18-DEC-2001; 2001US-0341189P.
XX 28-FEB-2002; 2002US-0361186P.
XX 12-MAR-2002; 2002US-0363673P.
XX 12-MAR-2002; 2002US-0363676P.
XX 03-JUN-2002; 2002US-00363676.
XX (CURA-) CURAGEN CORP.
XX
XX Anderson DW, Baumgartner JC, Boldog FI, Casman SJ, Edinger SR;
XX Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalt T, Kekuda R, Li L;
XX MacDougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
XX Pena CEA, Rastelli L, Shimkets RA, Stone DJ, Spytek KA, Vernet CAM;
XX Voss EZ, Zerhusen BD;
XX
XX WPI; 2003-140627/13.
XX N-PSDB; ACD03647.
XX

XX New NOVX polypeptides and nucleic acids, useful for preventing or
XX treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
XX atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
XX pharmacogenomics.
XX
XX Claim 1; Page 143; 332pp; English.

XX The invention describes an isolated polypeptide (I) comprising any of 27
XX 118-961 residue amino acid sequences, given in the specification, a
XX mature form of them, a sequence that is at least 95 % identical to them,
XX or a sequence having one or more conservative substitutions in them. The
XX polypeptide is useful in manufacturing a medicament for treating a
XX syndrome associated with a human disease selected from a pathology
XX associated with the polypeptide. The NOVX polypeptides, polynucleotides
XX and antibodies are useful in treating or preventing NOVX-associated
XX disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
XX disorders, AIDS, obesity, asthma, hematopoietic disorders, Parkinson's
XX disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
XX associated cachexia, and other wasting disorders associated with chronic
XX diseases. The nucleic acids and polypeptides may also be used as targets
XX for the identification of small molecules that modulate or inhibit e.g.
XX neurogenesis, cell differentiation, cell proliferation, hematopoiesis,
XX wound healing and angiogenesis. In gene therapy, in generation of
XX antibodies that bind immunospecifically to NOVX substances for use in
XX therapeutic or diagnostic methods. The nucleic acids are further used as
XX hybridisation probes, in chromosome mapping, tissue typing, preventive
XX medicine, and pharmacogenomics. The polypeptides are also useful as

CC vaccines. This is the amino acid sequence of a novel human G-protein
CC coupled receptor related protein NOV
XX
XX Sequence 398 AA;
SQ
Query Match 90.0%; Score 618; DB 6; Length 398;
Best Local Similarity 100.0%; Pred. No. 2.8e-62;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GKDFVQPTKICVGCPRDIPINSPELETHTITKLNAENNATFYFKIDNVKKARQV 62
DB 224 GKDFVQPTKICVGCPRDIPINSPELETHTITKLNAENNATFYFKIDNVKKARQV 283
QY 63 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTV 118
DB 284 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTV 339

RESULT 8
ADE76864
ID ADE76864 standard; protein; 427 AA.
XX
XX ADE76864;
XX
XX 29-JAN-2004 (first entry)
XX Human protein expressed in a liver disorder #9.
XX human; liver disorder; hyperlipidaemia; hypertension; type II diabetes;
KW tumour; liver; inflammatory disorder; immune response disorder;
KW high-throughput screening; differential gene expression; gene therapy.
XX Homo sapiens.
XX US2003108871-A1.
XX 12-JUN-2003.
XX 30-JUL-2001; 2001US-00919039.
XX 28-JUL-2000; 2000US-0222113P.
XX (KASE/) KASER M R.
XX Kaser MR;
XX WPI; 2004-031227/03.
XX N-PSDB; ADE76863.
XX Composition comprising several cDNAs that are differentially expressed in
XX treated human C3A liver cell cultures, useful for treating liver
XX disorders.
XX Claim 1; SEQ ID NO 29; 41pp; English.

XX The invention relates to a composition comprising several cDNAs that are
XX differentially expressed in a liver disorder. The composition is useful
XX for treating liver disorder such as hyperlipidaemia, hypertension, type
XX II diabetes, tumours of the liver and disorders of the inflammatory and
XX immune response. The composition is useful for a high-throughput method
XX of screening several molecules or compounds to identify a ligand which
XX specifically binds a cDNA. A protein encoded by the cDNA is useful for a
XX high-throughput method for using a protein to screen several molecules or
XX compounds to identify at least one ligand which specifically binds the
XX protein which involves combining the protein encoded by the cDNA with
XX several of molecules or compounds under conditions to allow specific
XX binding, and detecting specific binding between the protein and a
XX molecule or compound, therefore identifying a ligand which specifically
XX binds the protein. The composition is useful for detecting and
XX quantifying differential gene expression, can be used in gene therapy, to
XX formulate prognosis and to design a treatment regimen and to monitor the
XX efficacy of treatment. The present sequence represents the amino acid
XX sequence of a protein encoded by a cDNA differentially expressed in a

CC liver disorder.
 XX Sequence 427 AA;
 SQ Query Match 90.0%; Score 618; DB 8; Length 427;
 Best Local Similarity 100.0%; Pred. No. 3.1e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPPKICVGCPRDPTNSPELEETLTHITIKLAENNAATFYFKIDNVKKARQVW 62
 DB 253 GKDFVQPPKICVGCPRDPTNSPELEETLTHITIKLAENNAATFYFKIDNVKKARQVW 312

QY 63 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTV 118
 DB 313 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTV 368

RESULT 9
 ABU99144
 ID ABU99144 standard; protein; 615 AA.
 AC ABU99144;
 XX
 DT 01-AUG-2003 (first entry)
 XX
 DE Novel human GPCR related protein NOV12b.
 XX
 KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiac; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 XX
 OS Homo sapiens.
 XX
 PN WO200299116-A2.
 XX
 PD 12-DEC-2002.
 XX
 PF 04-JUN-2002; 2002WO-US017428.
 XX
 PR 04-JUN-2001; 2001US-0295607P.
 PR 04-JUN-2001; 2001US-0295661P.
 PR 06-JUN-2001; 2001US-0296404P.
 PR 14-JUN-2001; 2001US-0296418P.
 PR 15-JUN-2001; 2001US-0298285P.
 PR 21-JUN-2001; 2001US-0298556P.
 PR 26-JUN-2001; 2001US-0299499P.
 PR 28-JUN-2001; 2001US-0300883P.
 PR 13-AUG-2001; 2001US-0311972P.
 PR 29-AUG-2001; 2001US-0315071P.
 PR 29-AUG-2001; 2001US-0315660P.
 PR 14-SEP-2001; 2001US-0322293P.
 PR 17-SEP-2001; 2001US-0322706P.
 PR 14-DEC-2001; 2001US-0341186P.
 PR 28-FEB-2002; 2002US-0361189P.
 PR 12-MAR-2002; 2002US-0363673P.
 PR 12-MAR-2002; 2002US-0363676P.
 PR 03-JUN-2002; 2002US-00363676.
 XX
 PA (CURA-) CURAGEN CORP.
 XX
 PI Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR;
 PI Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalt T, Kekuda R, Li L;
 PI MacDougall JR, Malyankar UM, Millet I, Padigar M, Patturajan M;
 PI Pena CEA, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet CAM;

PI Voss EZ, Zerhusen BD;
 XX WPI; 2003-140627/13.
 DR N-PSDB; ACD03648.
 XX
 PT New NOVX polypeptides and nucleic acids, useful for preventing or
 PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX
 PS Claim 1; Page 144; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (1) comprising any of 27
 CC 118-561 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX Sequence 615 AA;
 SQ Query Match 90.0%; Score 618; DB 6; Length 615;
 Best Local Similarity 100.0%; Pred. No. 5.1e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPPKICVGCPRDPTNSPELEETLTHITIKLAENNAATFYFKIDNVKKARQVW 62
 DB 224 GKDFVQPPKICVGCPRDPTNSPELEETLTHITIKLAENNAATFYFKIDNVKKARQVW 283

QY 63 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTV 118
 DB 284 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTV 339

RESULT 10
 ABB78707
 ID ABB78707 standard; protein; 626 AA.
 XX
 AC ABB78707;
 XX
 DT 18-JUL-2002 (first entry)
 XX
 DE Human high molecular weight kininogen (HK) mature protein SEQ ID NO:1.
 XX
 KW Human; kininogen; high molecular weight kininogen; HK; D5 domain;
 KW D5 receptor; angiogenesis; endothelial cell; cytosolic; antitumour;
 KW antiatherosclerotic; vasotropic; vulnerary; tranquilliser; thrombolytic;
 KW ophthalmological; gynaecological; antidiabetic; antihypertensive;
 KW antiangiogenic; antiapoptotic; endocrine; apoptosis; gene therapy.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT Domain 384..508
 FT /label= D5_domain
 XX
 PN WO200214369-A2.

XX 21-FEB-2002.
 XX 24-JUL-2001; 2001WO-US023185.
 XX 24-JUL-2000; 2000US-0220194P.
 XX (ATTE-) ATTENUON LLC.
 XX Mazar AP, Juarez JC;
 XX WPI; 2002-393611/42.
 XX Novel human kininogen D5 domain polypeptides useful for treating
 XX conditions associated with endothelial cell migration, proliferation,
 XX invasion or angiogenesis, e.g. arthritis, macular degeneration, benign
 XX hyperplasia.
 XX Disclosure; Page 13; 84pp; English.
 XX The present invention describes an isolated polypeptide (I) that
 XX corresponds to the D5 domain of human kininogen, or biologically active
 XX peptide fragment, homologue or functional derivative, and which: (a)
 XX inhibits angiogenesis; (b) binds to the D5 binding site on endothelial
 XX cells (EC); (c) activates signalling pathways leading to the introduction
 XX of apoptosis in EC; and/or (d) inhibits the signalling pathway required
 XX for maintenance of EC viability. (I) has cytostatic, antitumour,
 XX antiatherosclerotic, vasotropic, vulnery, tranquiliser, thrombolytic,
 XX ophthalmological, gynaecological, antiulcer, antidiabetic, antiarthritic,
 XX antiangiogenic, antiapoptotic and endocrine activities. An antibody (IX)
 XX specific for an epitope of (I) is useful for inhibiting tumour growth or
 XX angiogenesis in a subject. (I), a D5 fusion polypeptide (II) or a dimeric
 XX or trimeric fusion polypeptide (III) can be used for inhibiting EC
 XX migration, proliferation, invasion, or angiogenesis, or for inducing EC
 XX apoptosis. An angiogenic EC-targeting pharmaceutical composition (X)
 XX comprising (I), (II), or (III), can be used for treating a subject having
 XX a disease or condition associated with undesired EC migration,
 XX proliferation, invasion or angiogenesis. (I), (II), or (III) can be used
 XX for isolating a D5 domain binding molecule from a complex mixture and for
 XX isolating or enriching cells expressing D5 domain binding sites from a
 XX cell mixture. The present sequence represents the mature human high
 XX molecular weight kininogen (HK) protein, which is given in the
 XX exemplification of the present invention
 XX SQ Sequence 626 AA;
 Query Match 90.0%; Score 618; DB 5; Length 626;
 Best Local Similarity 100.0%; Pred. No. 5.3e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITTKLNAENNAFFYFKIDNVKKARVQV 62
 DB 235 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITTKLNAENNAFFYFKIDNVKKARVQV 294
 QY 63 AGKYFIDFVARETTCSKESNEELTESCTKKLGSLDCNAEVVVPWEKKIYPTV 118
 DB 295 AGKYFIDFVARETTCSKESNEELTESCTKKLGSLDCNAEVVVPWEKKIYPTV 350
 RESULT 11
 ABG21101
 ID ABG21101 standard; protein; 644 AA.
 XX AC ABG21101;
 XX 18-FEB-2002 (first entry)
 XX Novel human diagnostic protein #21092.
 XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
 XX food supplement; medical imaging; diagnostic; genetic disorder.
 XX Homo sapiens.

XX WO200175067-A2.
 XX 11-OCT-2001.
 XX 30-MAR-2001; 2001WO-US008631.
 XX 31-MAR-2000; 2000US-00540217.
 XX 23-AUG-2000; 2000US-00649167.
 XX (HYSE-) HYSEQ INC.
 XX Drmanac RT, Liu C, Tang YT;
 XX WPI; 2001-639362/73.
 XX N-PSDB; AAS85288.
 XX New isolated polynucleotide and encoded polypeptides, useful in
 XX diagnostics, forensics, gene mapping, identification of mutations
 XX responsible for genetic disorders or other traits and to assess
 XX biodiversity.
 XX Claim 20; SEQ ID NO 51460; 103pp; English.
 XX The invention relates to isolated polynucleotide (I) and polypeptide (II)
 XX sequences. (II) is useful as hybridisation probes, polymerase chain
 XX reaction (PCR) primers, oligomers, and for chromosome and gene mapping.
 XX and in recombinant production of (II). The polynucleotides are also used
 XX in diagnostics as expressed sequence tags for identifying expressed
 XX genes. (I) is useful in gene therapy techniques to restore normal
 XX activity of (II) or to treat disease states involving (II). (II) is
 XX useful for generating antibodies against it, detecting or quantitating a
 XX polypeptide in tissue, as molecular weight markers and as a food
 XX supplement. (II) and its binding partners are useful in medical imaging
 XX of sites expressing (II). (I) and (II) are useful for treating disorders
 XX involving aberrant protein expression or biological activities. The
 XX polypeptide and polynucleotide sequences have applications in
 XX diagnostics, forensics, gene mapping, identification of mutations
 XX responsible for genetic disorders or other traits to assess biodiversity
 XX and to produce other types of data and products dependent on DNA and
 XX amino acid sequences. ABG0010-ABG30377 represent novel human diagnostic
 XX amino acid sequences of the invention. Note: The sequence data for this
 XX patent did not appear in the printed specification, but was obtained in
 XX electronic format directly from WIPO at
 XX ftp.wipo.int/pub/published_pct_sequences
 XX SQ Sequence 644 AA;
 Query Match 90.0%; Score 618; DB 4; Length 644;
 Best Local Similarity 100.0%; Pred. No. 5.5e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITTKLNAENNAFFYFKIDNVKKARVQV 62
 DB 253 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITTKLNAENNAFFYFKIDNVKKARVQV 312
 QY 63 AGKYFIDFVARETTCSKESNEELTESCTKKLGSLDCNAEVVVPWEKKIYPTV 118
 DB 313 AGKYFIDFVARETTCSKESNEELTESCTKKLGSLDCNAEVVVPWEKKIYPTV 368
 RESULT 12
 ABB78710
 ID ABB78710 standard; protein; 644 AA.
 XX AC ABB78710;
 XX 18-JUL-2002 (first entry)
 XX Human high molecular weight kininogen (HK) protein.
 XX Human; kininogen; high molecular weight kininogen; HK; D5 domain;
 XX D5 receptor; angiogenesis; endothelial cell; cytostatic; antitumour;

KW antiatherosclerotic; vasotropic; vulnerary; tranquilliser; thrombolytic;
 KW ophthalmological; gynaecological; antidiabetic; antidiabetic; antidiabetic;
 KW antiangiogenic; antiapoptotic; endocrine; apoptosis; gene therapy.

OS Hemo sapiens.

XX Key Location/Qualifiers
 FH Peptide 1..18
 FT /label= signal
 FT Protein 19..644
 FT /label= mature_human_high_molecular_weight_kinogen
 FT Disulfide-bond 28..614
 FT Disulfide-bond 83..94
 FT Disulfide-bond 107..126
 FT Disulfide-bond 142..145
 FT Disulfide-bond 206..218
 FT Disulfide-bond 229..248
 FT Disulfide-bond 264..267
 FT Disulfide-bond 328..340
 FT Disulfide-bond 351..370
 FT Domain 402..526
 FT /label= D5_domain

WO200214369-A2.

21-FEB-2002.

24-JUL-2001; 2001WO-US023185.

24-JUL-2000; 2000US-0220194P.

(ATTE-) ATTENDON LLC.

Mazar AP, Juarez JC;

WPI; 2002-393611/42.

Novel human kininogen D5 domain polypeptides useful for treating conditions associated with endothelial cell migration, proliferation, invasion or angiogenesis, e.g. arthritis, macular degeneration, benign hyperplasia.

Disclosure; Fig 1B-E; 84pp; English.

The present invention describes an isolated polypeptide (I) that corresponds to the D5 domain of human kininogen, or biologically active peptide fragment, homologue or functional derivative, and which: (a) inhibits angiogenesis; (b) binds to the D5 binding site on endothelial cells (EC); (c) activates signalling pathways leading to the introduction of apoptosis in EC; and/or (d) inhibits the signalling pathway required for maintenance of EC viability. (I) has cytostatic, antitumour, antiatherosclerotic, vasotropic, vulnerary, tranquilliser, thrombolytic, ophthalmological, gynaecological, antidiabetic, antidiabetic, antidiabetic, antidiabetic, antidiabetic and endocrine activities. An antibody (IX) specific for an epitope of (I) is useful for inhibiting tumour growth or angiogenesis in a subject. (I), a D5 fusion polypeptide (II) or a dimeric or trimeric fusion polypeptide (III) can be used for inhibiting EC migration, proliferation, invasion, or angiogenesis, or for inducing EC apoptosis. An angiogenic EC-targeting pharmaceutical composition (X) comprising (I), (II), or (III), can be used for treating a subject having a disease or condition associated with undesired EC migration, proliferation, invasion or angiogenesis. (I), (II), or (III) can be used for isolating a D5 domain binding molecule from a complex mixture and for isolating or enriching cells expressing D5 domain binding sites from a cell mixture. The present sequence represents the human high molecular weight kininogen (HK) protein, which is given in the exemplification of the present invention

Sequence 644 AA;

Query Match 90.0%; Score 618; DB 5; Length 644;

Best Local Similarity 100.0%; Pred. No. 5.5e-62;

Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 GKDFVQPTKICVGCPRDPTNSPELEETLTHITIKLNAENATFYFKIDNVKKARQV 62
 Db 253 GKDFVQPTKICVGCPRDPTNSPELEETLTHITIKLNAENATFYFKIDNVKKARQV 312
 Qy 63 AGKKYFIDFVARETTCSKESNEELTESCTKKLGSLDCNAEVVVPWEKKIYPTV 118
 Db 313 AGKKYFIDFVARETTCSKESNEELTESCTKKLGSLDCNAEVVVPWEKKIYPTV 368

RESULT 13

ABU99150

ID ABU99150 standard; protein; 644 AA.

XX AC ABU99150;

XX DT 01-AUG-2003 (first entry)

XX DE Novel human GPCR related protein NOV12h.

XX KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiac; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.

XX OS Homo sapiens.

XX PN WO200299116-A2.

XX PD 12-DEC-2002.

XX PF 04-JUN-2002; 2002WO-US017428.

XX PR 04-JUN-2001; 2001US-0295607P.

XX PR 04-JUN-2001; 2001US-0295661P.

XX PR 06-JUN-2001; 2001US-0298404P.

XX PR 06-JUN-2001; 2001US-0298418P.

XX PR 14-JUN-2001; 2001US-0298285P.

XX PR 15-JUN-2001; 2001US-0298556P.

XX PR 21-JUN-2001; 2001US-0299949P.

XX PR 26-JUN-2001; 2001US-0300883P.

XX PR 28-JUN-2001; 2001US-0301550P.

XX PR 13-AUG-2001; 2001US-0311972P.

XX PR 27-AUG-2001; 2001US-0315071P.

XX PR 29-AUG-2001; 2001US-0315660P.

XX PR 14-SEP-2001; 2001US-0322293P.

XX PR 17-SEP-2001; 2001US-0322706P.

XX PR 14-DEC-2001; 2001US-0341186P.

XX PR 28-FEB-2002; 2002US-0361189P.

XX PR 12-MAR-2002; 2002US-0363673P.

XX PR 12-MAR-2002; 2002US-0363676P.

XX PR 03-JUN-2002; 2002US-00363676.

(CURA-) CURAGEN CORP.

XX Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR;

XX Gangolli EA, Gerlach VL, Gorman I, Guo X, Hjalt T, Kekuda R, Li L;

XX Macdougall JR, Malyankar UM, Millet I, Padigar M, Patturajan M; CAV;

XX Pena CE, Rastelli L, Shimkets RA, Stone DJ, Spytek KA, Vernet

XX Voss EZ, Zerhusen BD;

XX WPI; 2003-140627/13.

XX DR N-PSDB; ACD03654.

XX New NOVX polypeptides and nucleic acids, useful for preventing or

XX treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,

PT	atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or pharmacogenomics.	
XX		
PS	Claim 1; Page 148; 332pp; English.	
XX		
CC	The invention describes an isolated polypeptide (I) comprising any of 27 118-961 residue amino acid sequences, given in the specification, a mature form of them, a sequence that is at least 95 % identical to them, or a sequence having one or more conservative substitutions in them. The polypeptide is useful in manufacturing a medicament for treating a syndrome associated with a human disease selected from a pathology associated with the polypeptide. The NOVX polypeptides, polynucleotides and antibodies are useful in treating or preventing NOVX-associated disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune disease, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's disease, Alzheimer's disease, infections, multiple sclerosis, cancer-associated cachexia, and other wasting disorders associated with chronic diseases. The nucleic acids and polypeptides may also be used as targets for the identification of small molecules that modulate or inhibit e.g. neurogenesis, cell differentiation, cell proliferation, haematopoiesis, wound healing and angiogenesis, in gene therapy, in generation of antibodies that bind immunospecifically to NOVX substances for use in therapeutic or diagnostic methods. The nucleic acids are further used as hybridisation probes, in chromosome mapping, tissue typing, preventive medicine, and pharmacogenomics. The polypeptides are also useful as vaccines. This is the amino acid sequence of a novel human G-protein coupled receptor related protein NOV	
XX		
SQ	Sequence 644 AA;	
Query Match 90.0%; Score 618; DB 6; Length 644;		
Best Local Similarity 100.0%; Pred. No. 5.5e-62;		
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		
Qy	3 GKDFVQPTKICVGCPRDIPNSELTELTHTTTKLAENNAATFYKIDNVKARQVV 62	
Db	253 GKDFVQPTKICVGCPRDIPNSELTELTHTTTKLAENNAATFYKIDNVKARQVV 312	
Qy	63 AGKYFIDFVARETTCSEKSENEELTESCETKLGSLDCKNAEVVVPWEKKIYPTV 118	
Db	313 AGKYFIDFVARETTCSEKSENEELTESCETKLGSLDCKNAEVVVPWEKKIYPTV 369	
RESULT 14		
ABU99145		
ID	ABU99145 standard; protein; 644 AA.	
XX		
AC	ABU99145;	
XX		
DT	01-AUG-2003 (first entry)	
XX		
DE	Novel human GPCR related protein NOV12c.	
XX		
KW	Human; G-protein coupled receptor related protein; GPCR related protein; NOV; cytosolic; cardiant; antiarteriosclerotic; antidiabetic;	
KW	immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;	
KW	antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;	
KW	NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;	
KW	diabetes; immune disorder; AIDS; obesity; asthma;	
KW	haematopoietic disorder; Parkinson's disease; Alzheimer's disease;	
KW	infection; multiple sclerosis; cancer-associated cachexia;	
KW	wasting disorder; chronic disease; neurogenesis; cell differentiation;	
KW	cell proliferation; haematopoiesis; wound healing; angiogenesis;	
KW	chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.	
OS	Homo sapiens.	
XX		
FN	WO200299116-A2.	
XX		
PD	12-DEC-2002.	
XX		
PF	04-JUN-2002; 2002WO-US017428.	
XX		

Db 313 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVVWPWEKKIYPTV 368

RESULT 15

AAB37447
ID AAB37447 standard; protein; 122 AA.

XX
XX

AC AAB37447;

XX 21-FEB-2001 (first entry)

XX DT

XX XX

XX Human kininogen D3.

XX DE

XX Enzyme; legumain; endopeptidase; cystatin; human; kininogen.

XX KW

XX XX

XX Homo sapiens.

XX OS

XX WO2000064945-A1.

XX FN

XX 02-NOV-2000.

XX PD

XX 20-APR-2000; 2000WO-GB001571.

XX PF

XX 22-APR-1999; 99GB-00009133.

XX PR

XX (BABR-) ABRAHAM INST.

XX PA

XX Abrahamson M, Barrett AJ;

XX PI

XX WPI; 2000-687316/67.

XX DR

XX Inhibition of mammalian legumain or legumain-related endopeptidase by

PT cystatin involves interaction with second papain-non-reactive site of

PT cystatin.

XX XX

XX Disclosure; Fig 4; 45pp; English.

XX XX

CC The present invention relates to inhibition of the enzymatic activity of

CC legumain or a legumain-related endopeptidase by cystatin. The inhibition

CC involves an interaction between legumain and a papain-non-reactive site

CC of cystatin. Legumain (EC 3.4.22.34) is a cysteine endopeptidase, and

CC performs a protein-processing function. The present sequence is human

CC kininogen D3, which was used in the present invention. Kininogen is a

CC type 3 cystatin

XX XX

SQ Sequence 122 AA;

Query Match 85.3%; Score 586; DB 3; Length 122;
Best Local Similarity 100.0%; Pred. No. 2.7e-59;
Matches 110; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 PPTKICVCGPRDIPNPSPELETLTHTITKLNANNATFYFKIDNVKKARVQVVGKKYF 68

Db 1 PPTKICVCGPRDIPNPSPELETLTHTITKLNANNATFYFKIDNVKKARVQVVGKKYF 60

QY 69 IDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVVWPWEKKIYPTV 118

Db 61 IDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVVWPWEKKIYPTV 110

Search completed: September 24, 2004, 14:08:38
Job time : 52.308 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:07:01 ; Search time 14,732 Seconds
(without alignments)
445.051 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687

Sequence: 1 GSGKDFVQPTKICVGCPRD.....VPWEKKIYPTVTNVHCECF 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues

Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents, AA.*
1: /cgn2_6/ptodata/2/iaa/5A_COMB.pep.*
2: /cgn2_6/ptodata/2/iaa/5B_COMB.pep.*
3: /cgn2_6/ptodata/2/iaa/6A_COMB.pep.*
4: /cgn2_6/ptodata/2/iaa/6B_COMB.pep.*
5: /cgn2_6/ptodata/2/iaa/PCTUS_COMB.pep.*
6: /cgn2_6/ptodata/2/iaa/backfiles.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	558	81.2	117	1	US-08-193-114B-1
2	556.5	81.0	117	5	PCT-US92-06809-1
3	163.5	23.8	145	2	US-08-832-535-2
4	163.5	23.8	145	3	US-09-019-485-2
5	163.5	23.8	145	3	US-09-019-485-3
6	163.5	23.8	145	3	US-09-431-480-9
7	163.5	23.8	145	3	US-09-617-302-9
8	163.5	23.8	145	4	US-09-528-436B-2
9	163	23.7	178	2	US-08-791-522-1
10	163	23.7	178	3	US-09-314-777-1
11	138.5	20.2	121	4	US-09-775-932-14
12	138.5	20.2	128	4	US-09-775-932-12
13	138.5	20.2	149	2	US-08-461-030C-2
14	138.5	20.2	149	3	US-08-744-138-2
15	138.5	20.2	149	3	US-09-431-480-8
16	138.5	20.2	149	3	US-09-431-480-10
17	138.5	20.2	149	3	US-09-617-302-8
18	138.5	20.2	149	3	US-09-617-302-10
19	138.5	20.2	149	4	US-09-241-376-2
20	138.5	20.2	149	4	US-09-940-497-2
21	137.5	20.0	112	4	US-08-849-932-16
22	136.5	19.9	118	4	US-09-775-932-24
23	135.5	19.7	146	6	5432264-6
24	134	19.5	148	5	PCT-US95-07135-2
25	132.5	19.3	120	4	US-09-775-932-2
26	132.5	19.3	145	2	US-08-832-535-11
27	132.5	19.3	146	2	US-08-791-522-3

28	132.5	19.3	146	3	US-08-744-138-3	Sequence 3, Appli
29	132.5	19.3	146	3	US-09-019-485-4	Sequence 4, Appli
30	132.5	19.3	146	3	US-09-314-777-3	Sequence 3, Appli
31	132.5	19.3	146	3	US-09-431-480-6	Sequence 6, Appli
32	132.5	19.3	146	3	US-09-617-302-6	Sequence 6, Appli
33	132.5	19.3	146	4	US-09-241-376-3	Sequence 3, Appli
34	132.5	19.3	146	4	US-09-528-436B-3	Sequence 3, Appli
35	132.5	19.3	146	4	US-09-886-319A-47	Sequence 47, Appli
36	132.5	19.3	146	4	US-09-940-497-3	Sequence 3, Appli
37	132.5	19.3	146	4	US-09-976-594-37	Sequence 37, Appli
38	132.5	19.3	146	4	US-08-849-303-17	Sequence 17, Appli
39	132.5	19.3	146	5	PCT-US95-07135-9	Sequence 9, Appli
40	132	19.3	26	3	US-08-676-242-15	Sequence 15, Appli
41	131.5	19.1	382	4	US-09-599-360B-93	Sequence 93, Appli
42	130	18.9	127	4	US-08-849-303-19	Sequence 19, Appli
43	129.5	18.9	140	4	US-09-886-319A-46	Sequence 46, Appli
44	129.5	18.9	140	4	US-09-886-319A-48	Sequence 48, Appli
45	128	18.6	111	4	US-08-849-303-26	Sequence 26, Appli

ALIGNMENTS

RESULT 1
US-08-193-114B-1
; Sequence 1, Application US/08193114B
; Patent No. 5472945
; GENERAL INFORMATION:
; APPLICANT: Schmaier, Alvin H.
; APPLICANT: Jiang, Yongping
; TITLE OF INVENTION: Modulation of Blood
; TITLE OF INVENTION: Pressure and Inhibition of Platelet Activation
; TITLE OF INVENTION: with Kininogen Fragment
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Seidel, Gonda, Lavorgna &
; ADDRESSEE: Monaco, P.C.
; STREET: 1800 Two Penn Center Plaza
; CITY: Philadelphia
; STATE: Pennsylvania
; COUNTRY: U.S.A.
; ZIP: 19102
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 720 Kb
; COMPUTER: IBM PS/2
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: WordPerfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/193,114B
; FILING DATE: 9 February 1994
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: U.S. Application
; APPLICATION NUMBER: Serial No. 5472945 07/744,545
; FILING DATE: 13 August 1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Monaco, Daniel A.
; REGISTRATION NUMBER: 30,480
; REFERENCE/DOCKET NUMBER: 6056-137 CII
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (215) 568-8383
; TELEFAX: (215) 568-5549
; TELEX: No. 5472945e
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 117 amino acids
; TYPE: peptide
; TOPOLOGY: linear
US-08-193-114B-1

Query Match 81.2%; Score 558; DB 1; Length 117;
Best Local Similarity 100.0%; Pred. No. 2.3e-55;
Matches 105; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 14 CVCGRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARVQVWAGKYYFDVA 73
Db 1 CVCGRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARVQVWAGKYYFDVA 60
QY 74 RETTCKSNBELLTSCCTKLGQSLDCNAEYVVPWEKKIYPTV 118
Db 61 RETTCKSNBELLTSCCTKLGQSLDCNAEYVVPWEKKIYPTV 105

RESULT 2
PCT-US92-06809-1
; Sequence 1, Application PC/TUS9206809
; GENERAL INFORMATION:
; APPLICANT: Schmaier, Alvin H.
; TITLE OF INVENTION: Modulation of Blood
; TITLE OF INVENTION: Pressure by Altering Bradykinin Levels
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Temple University - Of the
; STREET: 406 University Services
; CITY: Philadelphia
; STATE: Pennsylvania
; COUNTRY: U.S.A.
; ZIP: 19122

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 720 Kb
; COMPUTER: IBM PS/2
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: WordPerfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US92/06809
; FILING DATE: 19910813
; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: U.S. Application
; APPLICATION NUMBER: Serial No. 744,545
; FILING DATE: 13 August 1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Monaco, Daniel A.
; REGISTRATION NUMBER: 30,480
; REFERENCE/DOCKET NUMBER: 6056-137
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (215) 568-8383
; TELEFAX: (215) 568-5549
; TELEX:

; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 117 amino acids
; TYPE: AMINO ACID
; TOPOLOGY: linear
PCT-US92-06809-1

Query Match 81.0%; Score 556.5; DB 5; Length 117;
Best Local Similarity 93.8%; Pred. No. 3.3e-55;
Matches 106; Conservative 1; Mismatches 1; Indels 5; Gaps 1;

QY 14 CVCGRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARVQVWAGKYYFDVA 73
Db 1 CVCGRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARVQVWAGKYYFDVA 60
QY 74 RETTCKSNBELLTSCCTKLGQSLDCNAEYVVPWEKKIYPTV 126
Db 61 RETTCKSNBELLTSCCTKLGQSLDCNAEYVVPWEKKIYPTV 108

RESULT 3
US-08-832-535-2
; Sequence 2, Application US/08832535
; Patent No. 5919658

; GENERAL INFORMATION:
; APPLICANT: NI, JIAN
; APPLICANT: LI, HAODONG
; APPLICANT: YU, GUO-LIANG
; APPLICANT: GENTZ, REINER L
; TITLE OF INVENTION: HUMAN CYSTATIN F
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HUMAN GENOME SCIENCES, INC.
; STREET: 9410 KEY WEST AVENUE
; CITY: ROCKVILLE
; STATE: MD
; COUNTRY: US
; ZIP: 20850

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/832,535
; FILING DATE: 03-APR-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: KIMBALL, PAUL C.
; REGISTRATION NUMBER: 34,610
; REFERENCE/DOCKET NUMBER: PF265
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (201) 994-1700
; TELEFAX: (201) 994-1744
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 145 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-832-535-2

Query Match 23.8%; Score 163.5; DB 2; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVGRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARVQVWAGKYYFD 70
Db 32 SRVPGPFKTKTNDGVLQAAARYSVEKFNNTNDMFLFKESRITRALVQIVKGLKMLE 91
QY 71 FVARETTCKSNBELLTSCCTKLGQSLDCNAEYVVPWEKKIYPTV 124
Db 92 VEIGRTTCKKNQHLRL-DDCDFTQHTLKTLSLCYSEVVWVFW-----LQHFE 138

RESULT 4
US-09-019-485-2
; Sequence 2, Application US/09019485
; Patent No. 6066617
; GENERAL INFORMATION:
; APPLICANT: Li, Haodong
; APPLICANT: Yu, Guo-liang
; APPLICANT: Gentz, Reiner
; APPLICANT: Ni, Jian
; TITLE OF INVENTION: Cystatin F
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: MD
; COUNTRY: US
; ZIP: 20850

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS

```
;
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/019,485
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Benson, Robert H.
; REGISTRATION NUMBER: 30,446
; REFERENCE/DOCKET NUMBER: PF265P1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 3013098504
; TELEFAX: 3013098439
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 145 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-09-019-485-2

Query Match 23.8%; Score 163.5; DB 3; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

;
; 11 TKICVCPREDIPTNSPELEETLTHITKLAENNAATFYFKIDNVKARVQVAGKYYFID 70
; 32 SRVXPGFPKTIKTNDPGVLQAAARYSVEKFNNCTNDMFLFKESRITRALVQIVKGLKYLE 91
;
; 71 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVYVWPWEKKIYPTVTNNHWE 124
; 92 VEIGRTTCKKNQHLRL-DDCDFQTNHTLKQTLSCYSEVVVWPW-----LQHFE 138
;
;
; RESULT 5
; US-09-019-485-3
; Sequence 3, Application US/09019485
; Patent No. 6066617
; GENERAL INFORMATION:
; APPLICANT: Li, Haodong
; APPLICANT: Yu, Guo-Liang
; APPLICANT: Gentz, Reiner
; APPLICANT: Ni, Jian
; TITLE OF INVENTION: Cystatin F
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: MD
; COUNTRY: US
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/019,485
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Benson, Robert H.
; REGISTRATION NUMBER: 30,446
; REFERENCE/DOCKET NUMBER: PF265P1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 3013098504
; TELEFAX: 3013098439
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 145 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
```

```
;
; MOLECULE TYPE: protein
; US-09-019-485-3

Query Match 23.8%; Score 163.5; DB 3; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

;
; 11 TKICVCPREDIPTNSPELEETLTHITKLAENNAATFYFKIDNVKARVQVAGKYYFID 70
; 32 SRVXPGFPKTIKTNDPGVLQAAARYSVEKFNNCTNDMFLFKESRITRALVQIVKGLKYLE 91
;
; 71 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVYVWPWEKKIYPTVTNNHWE 124
; 92 VEIGRTTCKKNQHLRL-DDCDFQTNHTLKQTLSCYSEVVVWPW-----LQHFE 138
;
;
; RESULT 6
; US-09-431-480-9
; Sequence 9, Application US/09431480
; Patent No. 6235708
; GENERAL INFORMATION:
; APPLICANT: Holloway, James L.
; APPLICANT: Feldhaus, Andrew
; TITLE OF INVENTION: TESTIS SPECIFIC CYSTATIN-LIKE PROTEIN CYSTATIN T
; FILE REFERENCE: 98-72
; CURRENT APPLICATION NUMBER: US/09/431,480
; EARLIER FILING DATE: 1999-11-01
; EARLIER APPLICATION NUMBER: 60/109,217
; EARLIER FILING DATE: 1998-11-20
; EARLIER APPLICATION NUMBER: 60/156,382
; EARLIER FILING DATE: 1999-09-28
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 9
; LENGTH: 145
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-431-480-9

Query Match 23.8%; Score 163.5; DB 3; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

;
; 11 TKICVCPREDIPTNSPELEETLTHITKLAENNAATFYFKIDNVKARVQVAGKYYFID 70
; 32 SRVXPGFPKTIKTNDPGVLQAAARYSVEKFNNCTNDMFLFKESRITRALVQIVKGLKYLE 91
;
; 71 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVYVWPWEKKIYPTVTNNHWE 124
; 92 VEIGRTTCKKNQHLRL-DDCDFQTNHTLKQTLSCYSEVVVWPW-----LQHFE 138
;
;
; RESULT 7
; US-09-617-302-9
; Sequence 9, Application US/09617302
; Patent No. 6245523
; GENERAL INFORMATION:
; APPLICANT: Holloway, James L.
; APPLICANT: Feldhaus, Andrew
; TITLE OF INVENTION: TESTIS SPECIFIC CYSTATIN-LIKE PROTEIN CYSTATIN T
; FILE REFERENCE: 98-72 C1
; CURRENT APPLICATION NUMBER: US/09/617,302
; CURRENT FILING DATE: 2000-07-17
; PRIOR APPLICATION NUMBER: 09/431,480
; PRIOR FILING DATE: 1999-11-01
; PRIOR APPLICATION NUMBER: 60/109,217
; PRIOR FILING DATE: 1998-11-20
; PRIOR APPLICATION NUMBER: 60/156,382
; PRIOR FILING DATE: 1999-09-28
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 9
; LENGTH: 145
```



```

; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 12
; LENGTH: 128
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-775-932-12

Query Match      20.2%; Score 138.5; DB 4; Length 128;
Best Local Similarity 31.5%; Pred. No. 5.2e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPNPSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWAGKYFI 69
DB 9 POERNVGLRDLSPDDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSLVAGIKYFL 68
QY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
DB 69 TWMGSTDCKRTRVTGDHVDLT-TCPLAAGAQOEKLCDFEVLVVPWQ 115

RESULT 13
US-08-461-030C-2
; Sequence 2, Application US/08461030C
; Patent No. 5985601
; GENERAL INFORMATION:
; APPLICANT: NI, Jian
; APPLICANT: Yu, Guo-Liang
; APPLICANT: Gentz, Reiner
; APPLICANT: Rosen, Craig A.
; TITLE OF INVENTION: Human Cystatin E
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Ave
; CITY: Rockville
; STATE: MD
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/461,030C
; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: A. Anders, Brookes
; REGISTRATION NUMBER: 36,373
; REFERENCE/DOCKET NUMBER: PF202
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 301-301-8504
; TELEFAX: 301-309-8439
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 149 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-461-030C-2

Query Match      20.2%; Score 138.5; DB 2; Length 149;
Best Local Similarity 31.5%; Pred. No. 6.4e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPNPSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWAGKYFI 69
DB 30 POERNVGLRDLSPDDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSLVAGIKYFL 89

; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 178 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; CLONE: 30443
; US-09-314-777-1

Query Match      23.7%; Score 163; DB 3; Length 178;
Best Local Similarity 34.0%; Pred. No. 1.4e-10;
Matches 35; Conservative 20; Mismatches 44; Indels 4; Gaps 2;

QY 11 TKICVGCPRDIPNPSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWAGKYFI 70
DB 54 SVKFGFPKTIKNDPGVLQAARYSVERNCTNDMLFKESRITRALVQIVGLKYMLE 113
QY 71 FVARETTCSKENBELTSCSE---TKLGLSDLCNAEVVVPW 110
DB 114 VEIGRTCKKNOHLRL-DCCDFQTNHTLQTLSCYSEVVVPW 155

RESULT 11
US-09-775-932-14;
; Sequence 14, Application US/09775932
; Patent No. 6534477
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 14
; LENGTH: 121
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-775-932-14

Query Match      20.2%; Score 138.5; DB 4; Length 121;
Best Local Similarity 31.5%; Pred. No. 4.9e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPNPSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWAGKYFI 69
DB 2 POERNVGLRDLSPDDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSLVAGIKYFL 61
QY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
DB 62 TWMGSTDCKRTRVTGDHVDLT-TCPLAAGAQOEKLCDFEVLVVPWQ 108

RESULT 12
US-09-775-932-12
; Sequence 12, Application US/09775932
; Patent No. 6534477
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05

```

```

; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 12
; LENGTH: 128
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-775-932-12

Query Match      20.2%; Score 138.5; DB 4; Length 128;
Best Local Similarity 31.5%; Pred. No. 5.2e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPNPSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWAGKYFI 69
DB 9 POERNVGLRDLSPDDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSLVAGIKYFL 68
QY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
DB 69 TWMGSTDCKRTRVTGDHVDLT-TCPLAAGAQOEKLCDFEVLVVPWQ 115

RESULT 13
US-08-461-030C-2
; Sequence 2, Application US/08461030C
; Patent No. 5985601
; GENERAL INFORMATION:
; APPLICANT: Ni, Jian
; APPLICANT: Yu, Guo-Liang
; APPLICANT: Gentz, Reiner
; APPLICANT: Rosen, Craig A.
; TITLE OF INVENTION: Human Cystatin E
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Ave
; CITY: Rockville
; STATE: MD
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/461,030C
; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: A. Anders, Brookes
; REGISTRATION NUMBER: 36,373
; REFERENCE/DOCKET NUMBER: PF202
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 301-301-8504
; TELEFAX: 301-309-8439
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 149 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-461-030C-2

Query Match      20.2%; Score 138.5; DB 2; Length 149;
Best Local Similarity 31.5%; Pred. No. 6.4e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPNPSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWAGKYFI 69
DB 30 POERNVGLRDLSPDDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSLVAGIKYFL 89

; TELCOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 178 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; CLONE: 30443
; US-09-314-777-1

Query Match      23.7%; Score 163; DB 3; Length 178;
Best Local Similarity 34.0%; Pred. No. 1.4e-10;
Matches 35; Conservative 20; Mismatches 44; Indels 4; Gaps 2;

QY 11 TKICVGCPRDIPNPSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWAGKYFI 70
DB 54 SVKFGFPKTIKNDPGVLQAAQYSEVFNNCTNDMLFKESRITRALVQIVGLKXYMLE 113
QY 71 FVARETTCSKENBELTSCSE---TKLGLQSLDCNAEVVVPW 110
DB 114 VEIGRTCKKNOHLRL-DCCDFQTNHTLKQTLSCYSEVVVPW 155

RESULT 11
US-09-775-932-14;
; Sequence 14, Application US/09775932
; Patent No. 6534477
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 14
; LENGTH: 121
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-775-932-14

Query Match      20.2%; Score 138.5; DB 4; Length 121;
Best Local Similarity 31.5%; Pred. No. 4.9e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPNPSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWAGKYFI 69
DB 2 POERNVGLRDLSPDDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSLVAGIKYFL 61
QY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
DB 62 TWMGSTDCKRTRVTGDHVDLT-TCPLAAGAQOEKLCDFEVLVVPWQ 108

RESULT 12
US-09-775-932-12
; Sequence 12, Application US/09775932
; Patent No. 6534477
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05

```

QY 70 DFVARETTCSKE----SNEELTESCETKKGQ--SLDCNAEYVVVPWE 111
Db 90 TMEMGSTDCRKTRVTGHDVLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 136

RESULT 14

US-08-744-138-2

; Sequence 2, Application US/08744138

; Patent No. 6011012

; GENERAL INFORMATION:

; APPLICANT: Gentz, Reiner L.

; APPLICANT: Ni, Jian

; APPLICANT: Rosen, Craig A.

; APPLICANT: Yu, Guo-Liang

; TITLE OF INVENTION: Human Cystatin E

; NUMBER OF SEQUENCES: 13

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Human Genome Sciences, Inc.

; STREET: 9410 Key West Avenue

; CITY: Rockville

; STATE: Maryland

; COUNTRY: USA

; ZIP: 20850

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent In Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/744,138

; FILING DATE:

; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:

; NAME: Brookes, A. Anders

; REGISTRATION NUMBER: 36,373

; REFERENCE/DOCKET NUMBER: PF202P1

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: 301 309 8504

; TELEFAX: 301 309 8512

; INFORMATION FOR SEQ ID NO: 2:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 149 amino acids

; TYPE: amino acid

; TOPOLOGY: linear

; MOLECULE TYPE: protein

; US-08-744-138-2

Query Match 20.2%; Score 138.5; DB 3; Length 149;
Best Local Similarity 31.5%; Pred. No. 6.4e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPITNSPELEETLTHITIKLNAENNATFYFKIDNVKKARVQVWVAGKKYFI 69

Db 30 PQERWVGELELDLSPDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSQLVAGIKYFL 89

QY 70 DFVARETTCSKE----SNEELTESCETKKGQ--SLDCNAEYVVVPWE 111

Db 90 TMEMGSTDCRKTRVTGHDVLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 136

RESULT 15

US-09-431-480-8

; Sequence 8, Application US/09431480

; Patent No. 6235708

; GENERAL INFORMATION:

; APPLICANT: Holloway, James L.

; APPLICANT: Feldhaus, Andrew

; TITLE OF INVENTION: TESTIS SPECIFIC CYSTATIN-LIKE PROTEIN CYSTATIN T

; FILE REFERENCE: 98-72

; CURRENT APPLICATION NUMBER: US/09/431,480

; CURRENT FILING DATE: 1999-11-01

; EARLIER APPLICATION NUMBER: 60/109,217

; EARLIER FILING DATE: 1998-11-20

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:05:18 ; Search time 36.576 Seconds
(without alignments)
1095.549 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687
Sequence: 1 GSKGDFQVPPTKICVGCPRD.....VPWEKKIYPTVTVNHWECEP 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL 25:*

1: sp_arChaea:*

2: sp_bacteria:*

3: sp_fungi:*

4: sp_human:*

5: sp_invertebrate:*

6: sp_mammal:*

7: sp_mhc:*

8: sp_organelle:*

9: sp_phages:*

10: sp_plant:*

11: sp_rodent:*

12: sp_virus:*

13: sp_vertebrate:*

14: sp_unclassified:*

15: sp_virus:*

16: sp_bacteriap:*

17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Description
1	383	55.7	140 6 Q7YRP6
2	381	55.5	423 11 P70517
3	378	55.0	430 11 Q63581
4	171.5	25.0	167 11 Q9QWU5
5	163.5	23.8	167 4 Q7Z4J8
6	152.5	22.2	462 13 Q7ZY91
7	152.5	22.2	462 13 Q7SYH2
8	152.5	22.2	465 13 Q801E5
9	131.5	19.1	140 11 Q9EPX9
10	123.5	18.0	455 13 Q800S8
11	119	17.3	388 11 Q8CB17
12	117.5	17.1	464 13 Q801Z5
13	113.5	16.5	148 5 Q9NH95
14	113	16.4	140 11 Q80Y72
15	111	16.2	146 11 Q8K397
16	111	16.2	149 11 Q9D1B1

17	108.5	15.8	112	13	Q98SR4	Q98SR4 acipenser s
18	108.5	15.8	112	13	Q98SR3	Q98SR3 acipenser s
19	107	15.6	139	5	Q9TY2	Q9TY2 caenorhabdi
20	106	15.4	300	13	Q801Z6	Q801Z6 cyprinus ca
21	105	15.3	109	5	Q9TY65	Q9TY65 onchocerca
22	105	15.3	127	5	P90698	P90698 brugia mala
23	104.5	15.2	149	11	Q8VHCL	Q8VHCL rattus norv
24	104.5	15.2	161	5	O16159	O16159 brugia mala
25	102.5	14.9	127	5	Q9U9A1	Q9U9A1 onchocerca
26	101	14.7	148	11	Q9JMH4	Q9JMH4 mus musculu
27	99	14.4	110	10	Q8SA65	Q8SA65 sandersonia
28	98.5	14.3	107	5	Q8TOY2	Q8TOY2 sarcophaga
29	98.5	14.3	125	5	Q25620	Q25620 onchocerca
30	97.5	14.2	134	10	O41825	O41825 zea mays (m
31	95.5	13.9	143	5	O61973	O61973 caenorhabdi
32	94	13.7	122	5	O44396	O44396 haemonchus
33	93	13.5	138	4	Q8WXU6	Q8WXU6 homo sapien
34	92	13.4	157	5	Q17108	Q17108 acanthochei
35	90	13.1	139	11	Q8K5A3	Q8K5A3 rattus norv
36	88.5	12.9	92	10	Q9FXN6	Q9FXN6 arabidopsis
37	88.5	12.9	116	10	Q8XS7	Q8XS7 arabidopsis
38	88.5	12.9	124	10	O41906	O41906 arabidopsis
39	88.5	12.9	125	10	O22202	O22202 arabidopsis
40	88.5	12.9	134	10	P93627	P93627 zea mays (m
41	88.5	12.9	134	10	Q41897	Q41897 zea mays (m
42	88	12.8	199	10	Q39270	Q39270 brassica ca
43	87.5	12.7	141	11	O9DAP1	O9DAP1 mus musculu
44	86.5	12.6	141	11	Q80ZNS	Q80ZNS mus musculu
45	86.5	12.6	349	6	O14502	O14502 cercopithec

ALIGNMENTS

RESULT 1

Q7YRP6 ID Q7YRP6 PRELIMINARY; PRT; 140 AA.

AC Q7YRP6; 01-OCT-2003 (TrEMBLrel. 25, Created)

DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)

DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)

DE Low molecular weight kininogen (Fragment).

GN KNG.

OS Sus scrofa (Pig).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.

OX NCBI_TaxID=9823;

RN [1]

RP SEQUENCE FROM N.A.

RA Vonnahme K.A., Fernando S.C., Ross J.A., Ashworth M.D., DeSilva U.,

RA Malayer J.R., Geisert R.D.;

RA "Porcine Endometrial and Conceptus Expression of Kininogens and Plasma

RT Kallikrein in Cyclic and Pregnant Gilts."

RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.

DR EMBL; AY321363; AAP85260.1; -

FT NON_TER

FT NON_TER

SQ SEQUENCE 140 AA; 15650 MW; 177837836603F777 CRC64;

Query Match 55.7%; Score 383; DB 6; Length 140;

Best Local Similarity 78.9%; Pred. No. 2.8e-30;

Matches 75; Conservative 5; Mismatches 15; Indels 0; Gaps 0;

Qy 24 NSPELEETLTHITTKLNAENNAATFYFKIDNVKARVQVAGKYFDIFVARETTCSKSN 83

Db 1 DSPDLEPLNHSIAKLNAENNAVFYFKIGVEKATVQVAGKYIVFTARETTCSKSN 60

Qy 84 EELTESCEETKLGSLDCNAEVVVPWEKKIYPTV 118

Db 61 EELTESCEIKKPGQILKCNASVVPWEKKIYPTV 95

RESULT 2

```

P70517
ID AC P70517 PRELIMINARY; PRT; 423 AA.
AC Q63581
DT 01-FEB-1997 (T-EMBLrel. 02, Created)
DT 01-FEB-1997 (T-EMBLrel. 02, Last sequence update)
DT 01-OCT-2003 (T-EMBLrel. 25, Last annotation update)
DE Major acute phase alpha-1 protein precursor (Fragment).
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RA Cole T.;
RL Submitted (OCT-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=85149311; PubMed=2579644;
RA Cole T., Inglis A., Nagashima M., Schreiber G.;
RT "Major acute-phase alpha(1)-protein in the rat: Structure, molecular
cloning, and regulation of mRNA levels.";
RL Biochem. Biophys. Res. Commun. 126:719-724(1985).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=85127561; PubMed=2578992;
RA Cole T., Inglis A.S., Roxburgh C.M., Howlett G.J., Schreiber G.;
RT "Major acute phase alpha1-protein of the rat is homologous to bovine
kininogen and contains the sequence for bradykinin: its synthesis is
regulated at the mRNA level.";
RL FEBS Lett. 182:57-61(1985).
DR EMBL; K02814; AAA41569.1; -.
DR PIR; S68034; S68034.
DR PIR; S68035; S68035.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; Cystatin; 3.
DR SMART; SM00043; CV; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Signal.
FT NON_TER 1 1
FT SIGNAL <1 11 POTENTIAL.
FT CHAIN 12 423 POTENTIAL.
FT CHAIN 371 379 POTENTIAL.
SQ SEQUENCE 423 AA; 46905 MW; F9B8BD3198547949 CRC64;

Query Match 55.5%; Score 381; DB 11; Length 423;
Best Local Similarity 62.1%; Pred. No. 1.6e-29;
Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNANNATFYFKIDNVKKARQVW 62
DB 245 GDDLPELLKPCRCGCPREIPVDSPELKEALGHSIARLNAQHNIHFYFKIDTVKKATQVW 304

QY 63 AGKKYFIDFVARETCSKESNEELTESCTKLGSLDCNAEIVVVPWEKKIYPTV 118
DB 305 AGVIYVIEFIARETCSKQSKTELTADCTKHLGSLNCNANVYRPNKWKVPTV 360

RESULT 3
ID Q63581 PRELIMINARY; PRT; 430 AA.
AC Q63581
DT 01-NOV-1996 (T-EMBLrel. 01, Created)
DT 01-NOV-1996 (T-EMBLrel. 01, Last sequence update)
DT 01-OCT-2003 (T-EMBLrel. 25, Last annotation update)
DE Rat T-kininogen (T-KG).
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90034172; PubMed=2806908;

```

```

RA Anderson K.P., Croyle M.L., Lingrel J.B.;
RT "Primary structure of a gene encoding rat T-kininogen.";
RL Gene 81:119-128(1989).
DR EMBL; M29080; AAA42251.1; -.
DR EMBL; M29083; AAA42251.1; JOINED.
DR EMBL; M29084; AAA42251.1; JOINED.
DR EMBL; M29091; AAA42251.1; JOINED.
DR EMBL; M29085; AAA42251.1; JOINED.
DR EMBL; M29086; AAA42251.1; JOINED.
DR EMBL; M29087; AAA42251.1; JOINED.
DR EMBL; M29088; AAA42251.1; JOINED.
DR EMBL; M29089; AAA42251.1; JOINED.
DR PIR; S68034; S68034.
DR PIR; S68035; S68035.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; Cystatin; 3.
DR SMART; SM00043; CV; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
SQ SEQUENCE 430 AA; 47618 MW; 4508DEF4BDC978C CRC64;

Query Match 55.0%; Score 378; DB 11; Length 430;
Best Local Similarity 62.1%; Pred. No. 3.1e-29;
Matches 72; Conservative 13; Mismatches 31; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNANNATFYFKIDNVKKARQVW 62
DB 252 GDDLPELLKPCRCGCPREIPVDSPELKEALGHSIARLNAQHNIHFYFKIDTVKKATQVW 311

QY 63 AGKKYFIDFVARETCSKESNEELTESCTKLGSLDCNAEIVVVPWEKKIYPTV 118
DB 312 AGVIYVIEFIARETCSKQSKTELTADCTKHLGSLNCNANVYRPNKWKVPTV 367

RESULT 4
ID Q9QWL5 PRELIMINARY; PRT; 167 AA.
AC Q9QWL5
DT 01-MAY-2000 (T-EMBLrel. 13, Created)
DT 01-MAY-2000 (T-EMBLrel. 13, Last sequence update)
DT 01-JUN-2003 (T-EMBLrel. 24, Last annotation update)
DE Murine CMAP (CYSTATIN F) (LEUKOCYSTATIN).
GN MURINE CMAP OR CST7.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA Morita M., Arakawa H., Yoshiuchi N.;
RT "A novel cystatin-like metastasis associated gene.";
RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Embryo;
RX MEDLINE=21085660; PubMed=11217851;
RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yananaka I.,
RA Saito T., Okazaki Y., Gojohori T., Bono H., Kasukawa T., Saito R.,
RA Fiedschmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
RA Kuehl P., Lewis S., Matsuo Y., Nikaide I., Pesole G., Quackenbush J.,
RA Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamliya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,

```



```
RA Hayashizaki Y.;
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:685-690(2001);
DR EMBL; AB015224; BAA34940.1; -;
DR EMBL; AK004420; BAB23298.1; -;
DR HSSP; P01034; 1G96.
DR MGD; MGI:1298217; Cst7.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 2.
DR SEQUENCE 167 AA; 18847 MW; 61F776D8445095FE CRC64;

Query Match 25.0%; Score 171.5; DB 11; Length 167;
Best Local Similarity 35.5%; Pred. No. 3.4e-09;
Matches 39; Conservative 22; Mismatches 42; Indels 7; Gaps 3;

QY 4 KDFVQPTKICVCGPRDPTNSPELEETLTHITIKLAENNATFYFKIDNVKKARVQVVA 63
DB 50 KDLI---SSVKPGFPKTIETNPGVLKAARHSVEKFNCTNDIFLFKESHVSKALVQVVK 106
QY 64 GKYPIDFVARETTCKESNEELTESCE---TKLGGSLDCNAEYVVPW 110
DB 107 GLKTMLEVKIGRTTCRKTMHQL-DNCDFOINPALKXETLYCYSEVWIPW 155

RESULT 5
Q724J8 PRELIMINARY; PRT; 167 AA.
AC Q724J8;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cystatin F (Leukocystatin).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Kainine N., Chen X., Ralfs A., Halleck A., Hines L., Eisenstein S.,
RA Koundinya M., Raphael J., Moreira D., Kelley T., LaBaer J., Lin Y.,
RA Phelan M., Farmer A.;
RL Submitted (AUG-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BT009825; AAP8827.1; -;
SQ SEQUENCE 167 AA; 18857 MW; E339025ASBD50177 CRC64;

Query Match 23.8%; Score 163.5; DB 4; Length 167;
Best Local Similarity 31.6%; Pred. No. 2.1e-08;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVCGPRDPTNSPELEETLTHITIKLAENNATFYFKIDNVKKARVQVVAQKYYFD 70
DB 54 SRVKFGFPKTIKTNDPGVLOARYSVEKFNCTNDIMFLFKESRITRALVQIVKGLKYLE 113
QY 71 FVARETTCKESNEELTESCE---TKLGGSLDCNAEYVVPWPKIYPTVTVAHWE 124
DB 114 VEIGRTCKKQCHRL-DDCFOINHLKQILSCYSEVWVPW-----LQHPF 160

RESULT 6
Q7ZY91 PRELIMINARY; PRT; 462 AA.
AC Q7ZY91;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Similar to fetuin B.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;

RA SEQUENCE FROM N.A.
TISSUE=Embryo;
RA Klein S., Strausberg R.;
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC043891; AAH43891.1; -;
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; CY; 2.
DR SEQUENCE 462 AA; 53185 MW; D7BAD339961739FB CRC64;

Query Match 22.2%; Score 152.5; DB 13; Length 462;
Best Local Similarity 38.8%; Pred. No. 8.4e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

QY 10 PTKICVCGPRDPTNSPELEETLTHITIKLAENNATFYFKIDNVKKARVQVVA 65
DB 142 PGVILSTCP-DCTANEETPTITADTLIAEYKNSNNTFYFKIDHIERVRSQWVGP 200
QY 66 KYFIDFVARETTCKESNEELTESC 90
DB 201 SYFIQTIKETDCMKTQENWLSNC 225

RESULT 7
Q7SYH2 PRELIMINARY; PRT; 462 AA.
AC Q7SYH2;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cystatin domain fetuin-like protein.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
TISSUE=Ventral midgut;
RA Costa R.M.B., Mason J., Lee M., Amaya E., Zorn A.M.;
RA "Novel gene expression domains reveal early patterning of the Xenopus
RT endoderm.";
RL Submitted (MAR-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY260732; AAP82289.1; -;
SQ SEQUENCE 462 AA; 53186 MW; 796F92774CC27721 CRC64;

Query Match 22.2%; Score 152.5; DB 13; Length 462;
Best Local Similarity 38.8%; Pred. No. 8.4e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

QY 10 PTKICVCGPRDPTNSPELEETLTHITIKLAENNATFYFKIDNVKKARVQVVA 65
DB 142 PGVILSTCP-DCTANEETPTITADTLIAEYKNSNNTFYFKIDHIERVRSQWVGP 200
QY 66 KYFIDFVARETTCKESNEELTESC 90
DB 201 SYFIQTIKETDCMKTQENWLSNC 225

RESULT 8
Q801E5 PRELIMINARY; PRT; 465 AA.
AC Q801E5;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical histidine-rich protein (Fragment).
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae;
OC Xenopodinae; Xenopus.
```

```

OY NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22480013; PubMed=12591597;
RA Chen Y., Jurgens K., Hollemann T., Claussen M., Ramadori G.,
RA Pieler T.;
RT "Cell-autonomous and signal-dependent expression of liver and
RT intestine marker genes in pluripotent precursor cells from Xenopus
RT embryos.";
RL Mech. Dev. 120:277-288(2003).
DR EMBL; AY186284; AAC31610.1; -.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; CY; 2.
DR Hypothetical protein.
KW NON_TER
FT 1
SQ SEQUENCE 465 AA; 53528 MW; 0B403AB4F78BFD4 CRC64;

Query Match 22.2%; Score 152.5; DB 13; Length 465;
Best Local Similarity 38.8%; Pred. No. 8.5e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

OY 10 PTKICVGPDRIPNTSPLEELTHT---ITKLAENNAATFYFKIDNVKKARQVWAGK 65
Db 145 PGVILSTCP-DCPTANEETITPTTETAEITIAEYNKDSNNTRYFKIDHIERVRSQWVGP 203

OY 66 KYFIDFVARETTCKESNEELTESC 90
Db 204 SYFIQFTKETDCMTQENVLSNC 228

RESULT 9
OY Q9EPX9 PRELIMINARY; PRT; 140 AA.
ID Q9EPX9
AC Q9EPX9;
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Cystatin C.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=BALB/c;
RX MEDLINE=21010502; PubMed=11144350;
RA Taupin P.J., Ray J., Fischer W.H., Suhr S.T., Hakansson K., Grubb A.,
RA Gage F.H.;
RT "FGF-2-Responsive neural stem cell proliferation requires Ccrg, a novel
RT autocrine/paracrine cofactor.";
RL Neuron 28:385-397(2000).
DR EMBL; AF311741; AAG40283.1; -.
DR HSP; P01034; IG96.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00287; CYSTATIN; 1.
FT CHAIN 21 140 CYSTATIN C.
FT VARIANT 16 16 A -> G.
FT VARIANT 84 84 L -> P.
FT VARIANT 84 84 L -> P.
SQ SEQUENCE 140 AA; 15517 MW; 3A563406DD58D785 CRC64;

Query Match 19.1%; Score 131.5; DB 11; Length 140;
Best Local Similarity 27.8%; Pred. No. 2.6e-05;
Matches 32; Conservative 26; Mismatches 48; Indels 9; Gaps 4;

OY 15 VCCPRDIPNTSPLEELTHTITKLAENNAATFYFKIDNVKKARQVWAGKYFIDFVAR 74
Db 30 LGAPEADANEGVRALDFAVSEYKNGSDNAYHSRAQVVRARKQLVAGVNYFLDVEMG 89

```

```

OY 75 ETTCSKESNEELTESC---ETKLGSLDCNRAEVVVPWPKKIYPTVTNNHCE 126
Db 90 RTTCTK-SQTNLTD-CFFHDQPHLMRKALCSFQIYSPWK---GPHSLTNFSCK 138

RESULT 10
OY Q800S8 PRELIMINARY; PRT; 455 AA.
ID Q800S8
AC Q800S8;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Fetuin-A.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RX TISSUE=Liver;
RA Jia F.;
RT "Danio rerio fetuin-A.";
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY217758; AAC61483.1; -.
DR GO; GO:0005874; C:microtubule; IEA.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR GO; GO:0005198; F:structural molecule activity; IEA.
DR GO; GO:0007018; F:microtubule-based movement; IEA.
DR InterPro; IPR002453; Beta_tubulin.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00228; TUBULIN B AUTOREG; 1.
DR PROSITE; PS00228; TUBULIN B AUTOREG; 1.
SQ SEQUENCE 455 AA; 50627 MW; D8228729268A2ACB CRC64;

Query Match 18.0%; Score 123.5; DB 13; Length 455;
Best Local Similarity 26.7%; Pred. No. 0.00063;
Matches 32; Conservative 23; Mismatches 46; Indels 19; Gaps 4;

OY 2 SGKDFVQPTKICVGPDRIPNTSPLEELTHTITKLAENNAATFYFKIDNVKKARQV- 60
Db 134 SHEDLV---KKPCDGLPLHEPKALSVNAALAKFNKSNHKSIFYFKLMVGRISOW 189

OY 61 VWAGKYFIDFVARETTCKESNEELTES------CETKLG-QSLDCNRAEVY 106
Db 190 MPMGQSYFTQFAIMETNCTKDPQNPQPEACKALCGDQATYGFCKSKVSGSEPEVECEIY 249

RESULT 11
OY Q8CB17 PRELIMINARY; PRT; 388 AA.
ID Q8CB17
AC Q8CB17;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Fetuin beta.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=C57BL/60; TISSUE=Vagina;
RX MEDLINE=22354683; PubMed=12466851;
RA The FANTOM Consortium.
RA the RIKEN Genome Exploration Research Group Phase I & II Team;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs.";
RL Nature 420:563-573(2002).
DR EMBL; AK037043; BAC29682.1; -.
DR MGD; MGI:1890221; Fetub.

```


GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:04:32 ; Search time 8.636 Seconds
(without alignments)
765.738 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 697
Sequence: 1 GSGKDFVQPTKICVGCPRD.....VPWEKKIYPTVTVNHNECF 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_42:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	618	90.0	644	1 KNG HUMAN	P01042 homo sapien
2	440	64.0	436	1 KNL1 BOVIN	P01046 bos taurus
3	440	64.0	621	1 KXH1 BOVIN	P01044 bos taurus
4	413	60.1	434	1 KNL2 BOVIN	P01047 bos taurus
5	413	60.1	619	1 KXH2 BOVIN	P01045 bos taurus
6	413	60.1	661	1 KNG MOUSE	O08677 mus musculus
7	410	59.7	639	1 KNG RAT	P08934 rattus norv
8	388	56.5	430	1 KNT2 RAT	P08932 rattus norv
9	380	55.3	430	1 KNT1 RAT	P01048 rattus norv
10	171.5	25.0	144	1 CYTF MOUSE	O09098 mus musculus
11	163.5	23.8	145	1 CYTF HUMAN	O76096 homo sapien
12	138.5	20.2	146	1 CYTC MACMU	O19092 macaca mula
13	138.5	20.2	149	1 CYTM HUMAN	O15828 homo sapien
14	137.5	20.0	148	1 CYTC BOVIN	P01035 bos taurus
15	135	19.7	378	1 FETB RAT	O9qx79 rattus norv
16	132.5	19.3	146	1 CYTC HUMAN	P01034 homo sapien
17	132.5	19.3	146	1 CYTC SAISC	O19093 salmire sci
18	131.5	19.1	382	1 FETB HUMAN	Q9ugm5 homo sapien
19	130	18.9	127	1 CYTC RAT	P14841 rattus norv
20	129.5	18.9	140	1 CYTC MOUSE	P21460 mus musculus
21	128	18.6	111	1 CYT BITAR	P08935 bitis ariet
22	124.5	18.1	141	1 CYTT HUMAN	P05228 homo sapien
23	124.5	18.1	148	1 CYTC RABIT	O97862 oryctolagus
24	122.5	17.8	116	1 CYT COTJA	P81061 coturnix co
25	119	17.3	388	1 FETB MOUSE	Q9gxcl mus musculus
26	118.5	17.2	139	1 CYT CHICK	P01038 gallus gall
27	113	16.4	141	1 CYTS RAT	P19113 rattus norv
28	109.5	15.9	141	1 CYTN HUMAN	P01037 homo sapien
29	108.5	15.8	141	1 CYTS HUMAN	P01036 homo sapien
30	107	15.6	130	1 CYT ONCKE	Q98967 oncorhynch
31	105.5	15.4	162	1 CYTX ONCVO	P22085 onchocerca
32	105	15.3	130	1 CYT ONCMY	Q91195 oncorhynch
33	104	15.1	129	1 CYT_CYPEA	P35481 cyprinus ca

ALIGNMENTS

RESULT 1
KNG HUMAN
ID KNG HUMAN STANDARD; PRT; 644 AA.
AC P01042; P01043;
DT 21-JUL-1986 (Rel. 01, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Kinogen precursor (Alpha-2-thiol proteinase inhibitor) [Contains:
DE Bradykinin].
GN KNG.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORMS HMW AND LMW).
RC TISSUE=Liver;
RX MEDLINE=85234582; PubMed=2989293;
RA Takagaki Y., Kitamura N., Nakanishi S.;
RT "Cloning and sequence analysis of cDNAs for human high molecular
RT weight and low molecular weight prekininogens. Primary structures of
RT two human prekininogens.";
RL J. Biol. Chem. 260:8601-8609(1985).
RN [2]
RP GENE STRUCTURE
RX MEDLINE=85234583; PubMed=2989294;
RA Kitamura N., Kitagawa H., Fukushima D., Takagaki Y., Miyata T.,
RA Nakanishi S.;
RT "Structural organization of the human kininogen gene and a model for
RT its evolution.";
RL J. Biol. Chem. 260:8610-8617(1985).
RN [3]
RP SEQUENCE OF 1-401 FROM N.A.
RX MEDLINE=85122621; PubMed=6411591;
RA Okubo I., Kurachi K., Takasawa T., Shiohara H., Sasaki M.;
RT "Isolation of a human cDNA for alpha 2-thiol proteinase inhibitor and
RT its identity with low molecular weight kininogen.";
RL Biochemistry 23:5691-5697(1984).
RN [4]
RP SEQUENCE OF 379-644.
RX MEDLINE=86030270; PubMed=4054110;
RA Lottspeich F., Kellermann J., Henschel A., Foertsch B.,
RA Mueller-Esterl W.;
RT "The amino acid sequence of the light chain of human high-molecular-
RT mass kininogen.";
RL Eur. J. Biochem. 152:307-314(1985).
RN [5]
RP SEQUENCE OF 381-389.
RX MEDLINE=90255622; PubMed=4952632;
RA Pierce J.V.;
RT "Structural features of plasma kinins and kininogens.";
RL Fed. Proc. 27:52-57(1968).
RN [6]
RP DISULFIDE BONDS.
RA Sueyoshi T., Miyata T., Kato H., Iwanaga S.;
RT "Disulfide bonds in bovine HMW kininogens.";

P31727 sarcophaga
Q94269 mus musculus
Q9h114 homo sapien
P32766 mus musculus
P29699 mus musculus
O60576 homo sapien
P81714 najia atra (
Q9h112 homo sapien
P28325 homo sapien
P31726 zea mays (m
P24090 rattus norv
P02765 homo sapien

34 102 14.8 122 1 CYTA SARPE
35 101 14.7 139 1 CS11 MOUSE
36 95 13.8 165 1 CSTL HUMAN
37 94.5 13.8 142 1 CST8 MOUSE
38 94.5 13.8 345 1 A2HS MOUSE
39 94 13.7 142 1 CST8 HUMAN
40 93 13.5 99 1 CYT NAJAT
41 93 13.5 137 1 CS11 HUMAN
42 93 13.5 142 1 CYTD HUMAN
43 91.5 13.3 135 1 CYTI NAIZE
44 91.5 13.3 352 1 A2HS RAT
45 88 12.8 367 1 A2HS HUMAN

RL Seikagaku 56:808-808(1984).
 [7]
 RN CARBOHYDRATE-LINKAGE SITE ASN-294.
 RX MEDLINE-22660472; PubMed-12754519;
 RA Zhang H., Li X.-J., Martin D.B., Abersold R.;
 RT "Identification and quantification of N-linked glycoproteins using
 RT hydrazide chemistry, stable isotope labeling and mass spectrometry.";
 RL Nat. Biotechnol. 21:660-666(2003).
 CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 CC HMW-kininogen plays an important role in blood coagulation by
 CC helping to position optimally prekallikrein and factor XI next to
 CC factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
 CC induced aggregation of thrombocytes; (4) the active peptide
 CC bradykinin that is released from HMW-kininogen shows a variety of
 CC physiological effects: (4A) influence in smooth muscle
 CC contraction, (4B) induction of hypotension, (4C) natriuresis and
 CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
 CC mediator of inflammation and causes (4E1) increase in vascular
 CC permeability, (4E2) stimulation of nociceptors (4E3) release of
 CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
 CC a cardioprotective effect (directly via bradykinin action,
 CC indirectly via endothelium-derived relaxing factor action); (5)
 CC LMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
 CC kininogen is in contrast to HMW-kininogen not involved in blood
 CC clotting.
 CC -1- SUBCELLULAR LOCATION: Secreted.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=2;
 CC Name=HMW;
 CC IsoId=P01042-1; Sequence=Displayed;
 CC Name=LMW;
 CC IsoId=P01042-2; Sequence=VSP_001261, VSP_001262;
 CC -1- TISSUE SPECIFICITY: Plasma.
 CC -1- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 CC -1- SIMILARITY: Contains 3 cystatin-like domains.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; K02566; AAB35497.1; --
 CC EMBL; M11437; AAB59550.1; --
 CC EMBL; M11438; AAB59550.1; JOINED.
 CC EMBL; M11521; AAB59550.1; JOINED.
 CC EMBL; M11522; AAB59550.1; JOINED.
 CC EMBL; M11523; AAB59550.1; JOINED.
 CC EMBL; M11524; AAB59550.1; JOINED.
 CC EMBL; M11525; AAB59550.1; JOINED.
 CC EMBL; M11526; AAB59550.1; JOINED.
 CC EMBL; M11527; AAB59550.1; JOINED.
 CC EMBL; M11528; AAB59550.1; JOINED.
 CC EMBL; M11437; AAB59551.1; --
 CC EMBL; M11438; AAB59551.1; JOINED.
 CC EMBL; M11521; AAB59551.1; JOINED.
 CC EMBL; M11522; AAB59551.1; JOINED.
 CC EMBL; M11523; AAB59551.1; JOINED.
 CC EMBL; M11524; AAB59551.1; JOINED.
 CC EMBL; M11525; AAB59551.1; JOINED.
 CC EMBL; M11526; AAB59551.1; JOINED.
 CC EMBL; M11527; AAB59551.1; JOINED.
 CC EMBL; M11528; AAB59551.1; JOINED.
 CC PIR; A01279; KGHUL1.
 CC PIR; A01280; KGHUL1.
 CC SWISS-2DPAGE; P01042; HUMAN.
 CC Genew; HGNC:6383; KNG.
 CC MIM; 228960; --
 CC GO; GO:0007596; P:blood coagulation; NAS.
 CC GO; GO:0030146; P:diuresis; NAS.
 CC GO; GO:0006954; P:inflammatory response; NAS.

DR GO; GO:0030147; P:natriuresis; NAS.
 DR GO; GO:0006939; P:smooth muscle contraction; NAS.
 DR InterPro; IPR000010; Cystatin.
 DR InterPro; IPR002395; Kininogen.
 DR Pfam; PF00031; cystatin; 3.
 DR PRINTS; PR00334; KININOGEN.
 DR SMART; SM00043; CY; 3.
 DR PROSITE; PS00287; CYSSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Thiol protease inhibitor; Vasodilator;
 KW Bradykinin; Blood coagulation; Inflammatory response; Signal;
 KW Alternative splicing; Pyrrolidone carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 644 KININOGEN.
 FT CHAIN 19 380 KININOGEN HEAVY CHAIN.
 FT PEPTIDE 381 389 BRADYKININ.
 FT CHAIN 390 644 KININOGEN LIGHT CHAIN.
 FT DOMAIN 19 136 CYSTATIN-LIKE 1.
 FT DOMAIN 137 258 CYSTATIN-LIKE 2.
 FT DOMAIN 259 380 CYSTATIN-LIKE 3.
 FT DOMAIN 420 510 HIS-RICH
 FT (ASSOCIATED WITH CLOTTING ACTIVITY).
 FT REPEAT 420 449
 FT REPEAT 450 479
 FT REPEAT 480 510
 FT MOD_RES 19 19
 FT DISULFID 28 614
 FT DISULFID 83 94
 FT DISULFID 107 126
 FT DISULFID 142 145
 FT DISULFID 206 218
 FT DISULFID 229 248
 FT DISULFID 264 267
 FT DISULFID 328 340
 FT DISULFID 351 370
 FT CARBOHYD 48 48
 FT CARBOHYD 169 169
 FT CARBOHYD 205 205
 FT CARBOHYD 294 294
 FT CARBOHYD 401 401
 FT CARBOHYD 533 533
 FT CARBOHYD 542 542
 FT CARBOHYD 546 546
 FT CARBOHYD 557 557
 FT CARBOHYD 571 571
 FT CARBOHYD 577 577
 FT CARBOHYD 593 593
 FT CARBOHYD 628 628
 FT VARSPLIC 402 427
 FT VARSPLIC 428 644
 FT CONFLICT 593 593
 FT SEQUENCE 644 AA; 71945 MW; 3132B4CBAF8FB7E CRC64;
 FT VARSPLIC 402 427
 FT VARSPLIC 428 644
 FT CONFLICT 593 593
 FT SEQUENCE 644 AA; 71945 MW; 3132B4CBAF8FB7E CRC64;
 Query Match 90.0%; Score 618; DB 1; Length 644;
 Best Local Similarity 100.0%; Pred. No. 3.4e-51;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDIPNTSPPELETLTITIKLNAENNAFYFKIDNVKKARQVQV 62
 DB 253 GKDFVQPTKICVGCPRDIPNTSPPELETLTITIKLNAENNAFYFKIDNVKKARQVQV 312
 QY 63 AGKVFIDFVARETTCSKESNEELTESCTKLGSLDCNAEYVVPWEKKIYPTV 118
 DB 313 AGKVFIDFVARETTCSKESNEELTESCTKLGSLDCNAEYVVPWEKKIYPTV 368
 RESULT 2
 ID_KNLI_BOVIN STANDARD; PRT; 436 AA.
 AC P01046;
 DT 21-JUL-1986 (Rel. 01, Created)

DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Kininogen, LMW I precursor (Thiol proteinase inhibitor) (Contains:
 DE Bradykinin).
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OC NCB1_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=93117859; PubMed=6572010;
 RA Nawa H., Kitamura N., Hirose T., Asai M., Inayama S., Nakanishi S.;
 RT "Primary structures of bovine liver low molecular weight kininogen
 RT precursors and their two mRNAs.";
 RL Proc. Natl. Acad. Sci. U.S.A. 80:90-94(1983).
 RN [2]
 RP SEQUENCE OF 19-378.
 RX MEDLINE=87137530; PubMed=3546295;
 RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
 RA Miyata T., Iwanaga S.;
 RT "Bovine high molecular weight kininogen. The amino acid sequence,
 RT positions of carbohydrate chains and disulfide bridges in the heavy
 RT chain portion.";
 RL J. Biol. Chem. 262:2768-2779(1987).
 CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 CC LMW-kininogen inhibits the aggregation of thrombocytes; (3) the
 CC active peptide kallidin that is released from LMW-kininogen shows
 CC a variety of physiological effects: (3A) influence in smooth
 CC muscle contraction, (3B) induction of hypotension, (3C)
 CC natriuresis and diuresis (kidney).
 CC -!- SUBCELLULAR LOCATION: Extracellular.
 CC -!- ALTERNATIVE PRODUCTS:
 CC Event-Alternative splicing; Named isoforms=2;
 CC Name=LMW I;
 CC IsoId=P01046-1; Sequence=Displayed;
 CC Name=HMW I;
 CC IsoId=P01044-1; Sequence=External;
 CC -!- TISSUE SPECIFICITY: Plasma.
 CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 CC -!- MISCELLANEOUS: LMW-kininogen is in contrast to HMW-kininogen not
 CC involved in blood clotting.
 CC -!- SIMILARITY: Contains 3 cystatin-like domains.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See [http://www.isb-sib.ch/](http://www.isb-sib.ch/announce/)
 CC or send an email to license@isb-sib.ch).
 CC
 CC EMBL; V00426; CAA23709.1; ..
 DR PIR; A01283; KGB011.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; cystatin; 3.
 DR SMART; SM00043; CY; 3.
 DR PROSITE; PS00287; CYSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;
 KW Thiol protease inhibitor; Bradykinin; Signal;
 KW Pyrrolidone carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 436 KININOGEN, LMW I.
 FT CHAIN 19 378 HEAVY CHAIN.
 FT PEPTIDE 380 388 BRADYKININ.
 FT CHAIN 389 436 LIGHT CHAIN.
 FT DOMAIN 135 135 CYSTATIN-LIKE 1.
 FT DOMAIN 136 257 CYSTATIN-LIKE 2.
 FT DOMAIN 238 378 CYSTATIN-LIKE 3.
 FT MOD RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
 FT MOD RES 19 19
 FT CARBOHYD 57 87 N-LINKED (GLCNAC. . .).
 FT CARBOHYD 136 136 O-LINKED (PARTIAL).
 FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
 FT

FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
 FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
 FT DISULFID 27 406 INTERCHAIN.
 FT DISULFID 82 93
 FT DISULFID 106 123
 FT DISULFID 141 144
 FT DISULFID 205 217
 FT DISULFID 228 247
 FT DISULFID 263 266
 FT DISULFID 327 339
 FT DISULFID 350 369
 FT CONFLICT 295 295
 SQ SEQUENCE 436 AA; 48427 MW; F01F7EB6914BCE6C CRC64;
 A -> T (IN REF. 1; CAA23709).
 Query Match 64.0%; Score 440; DB 1; Length 436;
 Best Local Similarity 70.4%; Pred. No. 1.8e-34;
 Matches 81; Conservative 14; Mismatches 20; Indels 0; Gaps 0;
 QY 4 KDFVPPPKICVGCPRDPTNSPELEBEILTHITIKLNENNAFFYFKIDNVKARQVVA 63
 DB 253 KDFVPPPKICVGCPRDPTNSPELEBEILTHITIKLNENNAFFYFKIDNVKARQVVA 312
 QY 64 GKYYFDVARETTCKESNEELTESCETKLGQSLDCNAEYVYVWPEKKIYPTV 118
 DB 313 GLKYSIVFIARETTCKSGNEELTKSCENINHGQILHCDANVYVWPEKKIYPTV 367
 RESULT 3
 KNHL_BOVIN STANDARD; PRT; 621 AA.
 ID KNHL_BOVIN
 AC P01044;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Kininogen, HMW I precursor (Thiol proteinase inhibitor) (Contains:
 DE Bradykinin).
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OC NCB1_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=84014106; PubMed=6571699;
 RA Kitamura N., Takagaki Y., Furuto S., Tanaka T., Nawa H., Nakanishi S.;
 RT "A single gene for bovine high molecular weight and low molecular
 RT weight kininogens.";
 RL Nature 305:545-549(1983).
 RN [2]
 RP SEQUENCE OF 19-378.
 RX MEDLINE=87137530; PubMed=3546295;
 RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
 RA Miyata T., Iwanaga S.;
 RT "Bovine high molecular weight kininogen. The amino acid sequence,
 RT positions of carbohydrate chains and disulfide bridges in the heavy
 RT chain portion.";
 RL J. Biol. Chem. 262:2768-2779(1987).
 RN [3]
 RP SEQUENCE OF 378-393.
 RX MEDLINE=70180420; PubMed=4986212;
 RA Kato H., Nagasawa S., Suzuki T.;
 RT "Studies on the structure of bovine kininogen: cleavages of disulfide
 RT bonds and of methionyl bonds in kininogen-II.";
 RL J. Biochem. 67:313-323(1970).
 RN [4]
 RP SEQUENCE OF 458-498.
 RX MEDLINE=75170265; PubMed=1169237;
 RA Han Y.N., Komiya M., Iwanaga S., Suzuki T.;
 RT "Studies on the primary structure of bovine high-molecular-weight
 RT kininogen. Amino acid sequence of a fragment ('histidine-rich
 RT peptide') released by plasma kallikrein.";
 RL J. Biochem. 77:55-68(1975).
 CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)

HMW-kininogen plays an important role in blood coagulation by helping to position optimally prekallikrein and factor XI next to factor XII. (3) HMW-kininogen inhibits the thrombin- and plasmin-induced aggregation of thrombocytes; (4) the active peptide bradykinin that is released from HMW-kininogen shows a variety of physiological effects: (4A) influence in smooth muscle contraction, (4B) induction of hypotension, (4C) natriuresis and diuresis, (4D) decrease in blood glucose level. (4E) it is a mediator of inflammation and causes (4E1) increase in vascular permeability, (4E2) stimulation of nociceptors (4E3) release of other mediators of inflammation (e.g. prostaglandins), (4F) it has a cardioprotective effect (derived via bradykinin action, indirectly via endothelium-derived relaxing factor action).

-1- SUBCELLULAR LOCATION: Extracellular.

-1- ALTERNATIVE PRODUCTS:

Event=Alternative splicing; Named isoforms=2;

Name=HMW II;

Isoid=P01044-1; Sequence=Displayed;

Name=LMW I;

Isoid=P01046-1; Sequence=External;

-1- TISSUE SPECIFICITY: Plasma.

-1- PTM: Bradykinin is released from kininogen by plasma kallikrein.

-1- SIMILARITY: Contains 3 cystatin-like domains.

This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <http://www.isb-sib.ch/announce/> or send an email to license@isb-sib.ch).

EMBL; A01491; CAA24735.1; -

PIR; A01284; KGBOL1.

InterPro; IPR000010; Cystatin.

InterPro; IPR002395; Kininogen.

Pfam; PF00031; Cystatin; 3.

PRINTS; PR00334; KININOGEN.

SMART; SM00043; CY; 3.

PROSITE; PS00287; CYSTATIN; 2.

Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;

Thiol protease inhibitor; Bradykinin; Blood coagulation;

Inflammatory response; Signal; Pyrrolidone carboxylic acid.

SIGNAL 1 18 PROBABLE

CHAIN 19 621 KININOGEN, HMW I.

CHAIN 19 378 HEAVY CHAIN.

PEPTIDE 380 388 BRADYKININ.

CHAIN 389 621 LIGHT CHAIN.

DOMAIN 19 135 CYSTATIN-LIKE 1.

DOMAIN 136 257 CYSTATIN-LIKE 2.

DOMAIN 258 378 CYSTATIN-LIKE 3.

MCN RES 19 19 PYRROLIDONE CARBOXYLIC ACID.

CARBOHYD 87 87 N-LINKED (GLCNAC. . .).

CARBOHYD 136 136 O-LINKED (PARTIAL. . .).

CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).

CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).

CARBOHYD 204 204 N-LINKED (GLCNAC. . .).

DISULFID 27 591 INTERCHAIN.

DISULFID 82 93

DISULFID 106 125

DISULFID 141 144

DISULFID 205 217

DISULFID 228 247

DISULFID 263 266

DISULFID 327 339

DISULFID 350 369

SEQUENCE 621 AA; 68890 MW; D16850BEFE3C55CD CRC64;

Query Match 64.0%; Score 440; DB 1; Length 621;

Best Local Similarity 70.4%; Pred. No. 2.7e-34;

Matches 81; Conservative 14; Mismatches 20; Indels 0; Gaps 0;

4 KDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNANNATFFKIDNVKARQVVA 63

Db 253 KDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNANNATFFKIDNVKARQVVA 312

QY 64 GKXFDIFVARETTCCKESNEELTESCETKKLGSLDCNAEVYVPMWEKIVPTV 118

Db 313 GLKYSIVFIARETTCCKSGSNEELTKSCEINHGILHCDANVYVPMWEKIVPTV 367

RESULT 4

KML2_BOVIN STANDARD; PRT; 434 AA.

ID KML2_BOVIN AC P01047;

DT 21-JUL-1986 (Rel. 01, Created)

DT 21-JUL-1986 (Rel. 01, Last sequence update)

DT 15-MAR-2004 (Rel. 43, Last annotation update)

DE Kininogen, LMW II precursor (Thiol proteinase inhibitor) [Contains: Bradykinin].

OS Bos taurus (Bovine).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovidae; Bovinae; Bos.

OX NCBI_TaxID=9913;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=83117859; PubMed=6572010;

RA Nawa H., Kitamura N., Hirose T., Asai M., Inayama S., Nakanishi S.; Miyata T., Iwanaga S.;

RT "Primary structures of bovine liver low molecular weight kininogen precursors and their two mRNAs";

RL Proc. Natl. Acad. Sci. U.S.A. 80:90-94 (1983).

RN [2]

RP SEQUENCE OF 19-376.

RX MEDLINE=87137530; PubMed=3546295;

RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H., Miyata T., Iwanaga S.;

RT "Bovine high molecular weight kininogen. The amino acid sequence, positions of carbohydrate chains and disulfide bridges in the heavy chain portion.";

RL J. Biol. Chem. 262:2768-2779 (1987).

CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2) active peptide kallidin that is released from LMW-kininogen shows a variety of physiological effects: (3A) influence in smooth muscle contraction, (3B) induction of hypotension, (3C) natriuresis and diuresis (kidney).

CC -1- SUBCELLULAR LOCATION: Extracellular.

CC -1- ALTERNATIVE PRODUCTS:

Event=Alternative splicing; Named isoforms=2;

Name=LMW II;

Isoid=P01047-1; Sequence=Displayed;

Name=HMW II;

Isoid=P01045-1; Sequence=External;

-1- TISSUE SPECIFICITY: Plasma.

-1- PTM: Bradykinin is released from kininogen by plasma kallikrein.

-1- MISCELLANEOUS: LMW-kininogen is in contrast to HMW-kininogen not involved in blood clotting.

-1- SIMILARITY: Contains 3 cystatin-like domains.

This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <http://www.isb-sib.ch/announce/> or send an email to license@isb-sib.ch).

EMBL; V00427; CAA23710.1; -

PIR; A01284; KGBOL2.

HSSP; P01038; 1A90.

InterPro; IPR000010; Cystatin.

Pfam; PF00031; Cystatin; 3.

SMART; SM00043; CY; 3.

PROSITE; PS00287; CYSTATIN; 2.

Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;

KW Thiol protease inhibitor; Bradykinin; Signal;
 KW Pyrrolidone carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 434 KININOGEN, LMW II.
 FT PEPTIDE 19 376 HEAVY CHAIN.
 FT CHAIN 378 386 BRADYKININ.
 FT CHAIN 387 434 LIGHT CHAIN.
 FT DOMAIN 19 135 CYSTATIN-LIKE 1.
 FT DOMAIN 136 256 CYSTATIN-LIKE 2.
 FT DOMAIN 257 376 CYSTATIN-LIKE 3.
 FT MOD RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
 FT CARBOHYD 87 87 N-LINKED (GLCNAC. . .).
 FT CARBOHYD 136 136 O-LINKED (PARTIAL. . .).
 FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
 FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
 FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
 FT CARBOHYD 280 280 N-LINKED (GLCNAC. . .).
 FT DISULFID 27 404 INTERCHAIN.
 FT DISULFID 82 93
 FT DISULFID 106 125
 FT DISULFID 141 144
 FT DISULFID 205 217
 FT DISULFID 228 247
 FT DISULFID 261 264
 FT DISULFID 325 337
 FT DISULFID 348 367
 SQ SEQUENCE 434 AA; 48148 MW; 73A7079DE3E03430 CRC64;
 Query Match 60.1%; Score 413; DB 1; Length 434;
 Best Local Similarity 67.2%; Pred. No. 6.6e-32;
 Matches 78; Conservative 14; Mismatches 22; Indels 2; Gaps 1;
 QY 3 GKDFVQPTKLCVGPDPNPSPELEBTLHTITKLAENNAFYFKIDNVKKARQV 62
 DB 252 GEDFL--PPWVCVCPKPIPVDSPLDEALNHSIAKLAHEHGTFFPKIDTVKATQV 309
 QY 63 AGKYFIDFARETTCKSESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTV 118
 DB 310 GGLKYSIVFIARETTCKSGSNEELTKSCIEINHGQILHCDAVYVVPWEKKVPTV 365
 RESULT 5
 ID KXH2_BOVIN STANDARD; PRT; 619 AA.
 AC P01045;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Kininogen, HMW II precursor (thiol proteinase inhibitor) [Contains:
 DE Bradykinin].
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OX NCBI_TaxId=9913;
 RN [1]
 RN SEQUENCE FROM N.A.
 RX MEDLINE=84014106; PubMed=6571699;
 RA Kitamura N., Takagaki Y., Furuto S., Tanaka T., Nawa H., Nakanishi S.;
 RA "A single gene for bovine high molecular weight and low molecular
 RT weight kininogens.";
 RL Nature 305:545-549 (1983).
 RN [2]
 RN SEQUENCE OF 19-376.
 RP MEDLINE=87137530; PubMed=3546295;
 RX Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
 RA Miyata T., Iwagawa S.;
 RA "Bovine high molecular weight kininogen. The amino acid sequence,
 RT positions of carbohydrate chains and disulfide bridges in the heavy
 RT chain portion.";
 RL J. Biol. Chem. 262:2768-2779 (1987).
 RN [3]
 RP SEQUENCE OF 376-391.

RX MEDLINE=70180420; PubMed=4986212;
 RA Kato H., Nagasawa S., Suzuki T.;
 RT "Studies on the structure of bovine kininogen: cleavages of disulfide
 bonds and of methionyl bonds in kininogen-II.";
 RL J. Biochem. 67:313-323 (1970).
 RN [4]
 RN SEQUENCE OF 387-455.
 RX MEDLINE=76260155; PubMed=956151;
 RA Han Y.N., Kato H., Iwanaga S., Suzuki T.;
 RT "Primary structure of bovine plasma high-molecular-weight kininogen.
 RT The amino acid sequence of a glycopeptide portion (fragment 1)
 RT following the C-terminus of the bradykinin moiety.";
 RL J. Biochem. 79:1201-1222 (1976).
 RN [5]
 RN SEQUENCE OF 456-496.
 RX MEDLINE=75170265; PubMed=1169237;
 RA Han Y.N., Komiyama M., Iwanaga S., Suzuki T.;
 RT "Studies on the primary structure of bovine high-molecular-weight
 RT kininogen. Amino acid sequence of a fragment ('histidine-rich
 RT peptide') released by plasma kallikrein.";
 RL J. Biochem. 77:55-68 (1975).
 CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 CC HMW-kininogen plays an important role in blood coagulation by
 CC helping to position optimally prekallikrein and factor XI next to
 CC factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
 CC induced aggregation of thrombocytes; (4) the active peptide
 CC bradykinin that is released from HMW-kininogen shows a variety of
 CC physiological effects: (4A) influence in smooth muscle
 CC contraction, (4B) induction of hypotension, (4C) natriuresis and
 CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
 CC mediator of inflammation and causes (4E1) increase in vascular
 CC permeability, (4E2) stimulation of nociceptors (4E3) release of
 CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
 CC a cardioprotective effect (directly via bradykinin action,
 CC indirectly via endothelium-derived relaxing factor action).
 CC -1- SUBCELLULAR LOCATION: Extracellular.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=2;
 CC Name=HMW II;
 CC IsoId=P01045-1; Sequence=Displayed;
 CC Name=LMW II;
 CC IsoId=P01047-1; Sequence=External;
 CC -1- TISSUE SPECIFICITY: Plasma.
 CC -1- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 CC -1- SIMILARITY: Contains 3 cystatin-like domains.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to licenses@isb-sib.ch).
 CC -----
 CC EMBL: V01492; CAA24736.1; -.
 CC F01: A01282; KGB0H2.
 CC H01: P01038; I01038.
 CC H02: P01038; I01038.
 CC InterPro: IPR000010; Cystatin.
 CC InterPro: IPR002395; Kininogen.
 CC Pfam: PF00031; Cystatin; 3.
 CC PRINTS: PR00334; KININOGEN.
 CC SMART: SM00043; CY; 3.
 CC PROSITE: PS00287; CYSTATIN; 2. Vasodilator; Alternative splicing;
 KW Glycoprotein; Plasma; Repeat; Thiol protease inhibitor; Bradykinin; Blood coagulation; Signal;
 KW Inflammatory response; Pyrrolidone carboxylic acid.
 FT SIGNAL 1 18 KININOGEN, HMW II.
 FT CHAIN 19 619 HEAVY CHAIN.
 FT CHAIN 19 376 BRADYKININ.
 FT PEPTIDE 378 386 LIGHT CHAIN.
 FT CHAIN 387 619 CYSTATIN-LIKE 1.
 FT DOMAIN 19 135 CYSTATIN-LIKE 2.
 FT DOMAIN 136 256

```

FT DOMAIN 257 376 CYPSTATIN-LIKE 3.
FT MOD_RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
FT CARBOHYD 87 87 N-LINKED (GLCNAC. . .).
FT CARBOHYD 136 136 O-LINKED (PARTIAL).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
FT CARBOHYD 280 280 N-LINKED (GLCNAC. . .).
FT CARBOHYD 400 400 O-LINKED.
FT DISULFID 27 589 INTERCHAIN.
FT DISULFID 82 93
FT DISULFID 106 125
FT DISULFID 141 144
FT DISULFID 205 217
FT DISULFID 228 247
FT DISULFID 261 264
FT DISULFID 325 337
FT DISULFID 348 367
FT VARIANT 398 398 T -> P.
FT VARIANT 401 401 L -> V.
FT VARIANT 454 454 H -> K.
SQ SEQUENCE 619 AA; F04320A8EB0EE0DA CRC64;

Query Match 60.1%; Score 413; DB 1; Length 619;
Best Local Similarity 67.2%; Pred. No. 9.9e-32;
Matches 78; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

QY 3 GKDFVPTKICVGCPRDPTNSPELEETLTHITKLAENNATFVKIDNVKKARVQV 62
Db 252 GEDFL--PPMVCVCGPKPVPDPSDLEALNHSIAKNAEHDTGYFKIDTVKKATQVV 309
QY 63 AGKYPIDFVARETTCSEKNEBELTSCCKLQSLDCAEYVYVPEKKIYPTV 118
Db 310 GGLKYSIVFIARETTCSEKNEBELTSCBEINTHGQILCHDANVYVPEEKYPTV 365

RESULT 6
KING_MOUSE STANDARD; PRT; 661 AA.
AC O08677; O08676; Q91XK5;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Kininogen precursor [Contains: Bradykinin].
GN KING.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORMS HMW AND LMW).
RC STRAIN=C57BL/6 X CBA; TISSUE=Liver;
RX MEDLINE=97342556; PubMed=9199253;
RA Okazaki Y., Furuno M., Kasukawa T., Adachi J., Bono H., Kondo S.,
RA Nikaide I., Osato N., Saito R., Suzuki H., Yamanaka I., Kiyosawa H.,
RA Yagi K., Tomaru Y., Hasegawa Y., Nogami A., Schonbach C., Gojobori T.,
RA Baldarelli R., Hill D.P., Bult C., Hume D.A., Quackenbush J.,
RA Schriml L.M., Kapin A., Matsuda H., Batalov S., Beisel K.W.,
RA Blake J.A., Bradt D., Brusio V., Chothia C., Corbani L.E., Cousins S.,
RA Dalla E., Dragani T.A., Fletcher C.F., Forrest A., Frazer K.S.,
RA Gaasterland T., Gariboldi M., Gissi C., Godzik A., Gough J.,
RA Grimm S., Gustincich S., Hirokawa N., Jackson I.J., Jarvis E.D.,
RA Kanai A., Kawaji H., Kawasawa Y., Kedzierski R.M., King B.L.,
RA Kongsava A., Kurochkin I.V., Lee Y., Lenhard B., Lyons P.A.,
RA Maglott D.R., Maltais L., Marchionni L., McKenzie L., Miki H.,

```

```

RA Nagashima T., Numata K., Okido T., Pavan W.J., Pertea G., Pesole G.,
RA Petrovsky N., Pillai R., Pontius J.U., Qi D., Ramachandran S.,
RA Ravasi T., Reed J.C., Reed D.J., Reid J., Ring B.Z., Ringwald M.,
RA Sancelin A., Schneider C., Sempie C.A., Setou M., Shimada K.,
RA Sultana R., Takeraka Y., Taylor M.S., Teasdale R.D., Tomita M.,
RA Verardo R., Wagner L., Wahlstedt C., Wang Y., Watanabe Y., Wells C.,
RA Wilming L.G., Wynshaw-Boris A., Yanagisawa M., Yang I., Yang L.,
RA Yuan Z., Zavolan M., Zhu Y., Zimmer A., Carninci P., Hayatsu N.,
RA Hirozane-Kishikawa T., Konno H., Nakamura M., Sakazume N., Sato K.,
RA Shiraki T., Waki K., Kawai J., Aizawa K., Arakawa T., Fukuda S.,
RA Hara A., Hashizume W., Imotani K., Ishii Y., Itoh M., Kagawa I.,
RA Miyazaki A., Sakai K., Sasaki D., Shibata K., Shinagawa A.,
RA Yasunishi A., Yoshino M., Waterston R., Lander E.S., Rogers J.,
RA Birney E., Hayashizaki Y.,
RA "Analysis of the mouse transcriptome based on functional annotation of
RA 60,770 full-length cDNAs.";
RA Nature 420:563-573 (2002).
RN [3]
RP SEQUENCE FROM N.A. (ISOFORM LMW).
RC TISSUE=Liver;
RX MEDLINE=22388257; PubMed=12477932;
RA Srausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wegner L., Shennen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buettow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.F., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.A., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Richards S., Morley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Wozny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kettelman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalhus D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RA "Generation and initial analysis of more than 15,000 full-length
RA human and mouse cDNA sequences.";
RN Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
CC -I- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC HMW-kininogen plays an important role in blood coagulation by
CC helping to position optimally prekallikrein and factor XI next to
CC factor XII; (3) HMW-kininogen inhibits the thrombin and plasmin-
CC induced aggregation of thrombocytes; (4) the active peptide
CC bradykinin that is released from HMW-kininogen shows a variety of
CC physiological effects: (4A) influence in smooth muscle
CC contraction, (4B) induction of hypotension, (4C) natriuresis and
CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
CC mediator of inflammation and causes (4E1) increase in vascular
CC permeability, (4E2) stimulation of nociceptors (4E3) release of
CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
CC a cardioprotective effect (directly via bradykinin action,
CC indirectly via endothelium-derived relaxing factor action); (5)
CC LMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
CC kininogen is in contrast to HMW-kininogen not involved in blood
CC clotting (By similarity).
CC -I- SUBCELLULAR LOCATION: Secreted.
CC -I- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=HMW;
CC IsoId=O08677-1; Sequence=Displayed;
CC Name=LMW;
CC IsoId=O08677-2; Sequence=VSP_001263, VSP_001264;
CC -I- TISSUE SPECIFICITY: Plasma.
CC -I- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -I- SIMILARITY: Contains 3 cystatin-like domains.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way

```



```

DR EMBL; M11884; AAA41487.1; -.
DR EMBL; M14369; AAA41484.1; -.
DR EMBL; M14369; AAA41485.1; ALT_SEQ.
DR EMBL; M14455; AAA41482.1; -.
DR PIR; A25486; A25486.
DR PIR; A28055; A28055.
DR InterPro; IPR000010; Cystatin.
DR PRINTS; PR000334; KININOGEN.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Glycoprotein; Plasma; Repeat; Thiol protease inhibitor; Vasodilator;
KW Bradykinin; Blood coagulation; Inflammatory response; Signal;
KW Alternative splicing; Multigene family.
FT SIGNAL 1 18
FT CHAIN 19 639 KININOGEN.
FT CHAIN 19 380 KININOGEN HEAVY CHAIN.
FT PEPTIDE 381 389 BRADYKININ.
FT CHAIN 390 639 KININOGEN LIGHT CHAIN.
FT DOMAIN 19 136 CYSTATIN-LIKE 1.
FT DOMAIN 137 238 CYSTATIN-LIKE 2.
FT DOMAIN 239 380 CYSTATIN-LIKE 3.
FT DOMAIN 439 514 HIS-RICH.
FT DISULFID 28 609 INTERCHAIN (BY SIMILARITY).
FT DISULFID 83 94 BY SIMILARITY.
FT DISULFID 107 126 BY SIMILARITY.
FT DISULFID 142 145 BY SIMILARITY.
FT DISULFID 206 218 BY SIMILARITY.
FT DISULFID 229 248 BY SIMILARITY.
FT DISULFID 264 267 BY SIMILARITY.
FT DISULFID 328 340 BY SIMILARITY.
FT DISULFID 351 370 BY SIMILARITY.
FT CARBOHYD 82 82 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 127 127 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 169 169 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 205 205 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 294 294 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 529 529 N-LINKED (GLCNAC. .) (POTENTIAL).
FT VARSPPLIC 402 433 VSPSPYKARVDEPDPGNEQPIHGHGMLHAKQ -> RLINS
FT FT CSYKRLKAGAPAPRQAESTVTP (in isoform
FT FT LWM)
FT FT /FTID=VSP 001265.
FT FT Missing (in isoform LWM).
FT FT /FTID=VSP 001266.
FT FT E -> K (IN REF. 2).
FT SEQUENCE 639 AA; 70933 NW; D3172DF94FF56AF5 CRC64;
Query Match 59.7%; Score 410; DB 1; Length 639;
Best Local Similarity 66.4%; Pred. No. 2e-31; Indels 0; Gaps 0;
Matches 77; Conservative 13; Mismatches 26;
QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKNAENNAATFYFKIDNVKKEARVQV 62
Db 253 GDDLFELLPEDCPGCPNPVDSPELKEALGHSIAQLNAENHHTFYFKIDTVKXATQV 312
QY 63 AGKKYFIDFVARETCTSKESNEELTESCTKLGSLDCNAEVVVVPEKKIYTV 118
Db 313 AGTKYVIEFIARETCTSKESNAELTADCTKRLGOSLNCNANVYMRPENKVVPTV 368
RESULT 8
KNT2_RAT
ID KNT2_RAT STANDARD; PRT; 430 AA.
AC P08932;
DT 01-NOV-1988 (Rel. 09, Created)
DT 01-NOV-1988 (Rel. 09, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE T-kininogen II precursor (Major acute phase protein) (Alpha-1-MAP)
DE (Thiostatin) [Contains: T-kinin].
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

```

```

OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=86008264; PubMed=2413018;
RA Furoto-Kato S., Matsumoto A., Kitamura N., Nakanishi S.;
RT "Primary structures of the mRNAs encoding the rat precursors for
RT bradykinin and T-kinin. Structural relationship of kininogens with
RT major acute phase protein and alpha 1-cysteine proteinase
RT inhibitor.";
RL J. Biol. Chem. 260:12054-12059 (1985).
CC -I- FUNCTION: Kininogens are plasma glycoproteins with a number of
CC functions: (1) as precursor of the active peptide bradykinin they
CC effect smooth muscle contraction, induction of hypotension and
CC increase of vascular permeability. (2) They play a role in blood
CC coagulation by helping to position optimally prekallikrein and
CC factor XI next to factor XII. (3) They are inhibitor of thiol
CC proteases.
CC -I- SUBCELLULAR LOCATION: Extracellular.
CC -I- TISSUE SPECIFICITY: Plasma.
CC -I- INDUCTION: In response to an inflammatory stimulant. T-kininogen
CC II synthesis is induced and the plasma concentration of
CC T-kininogen I is raised.
CC -I- PTM: AS T-KININ IS PRECEDED BY A MET INSTEAD OF AN ARG OR LYS, IT
CC IS NOT RELEASED FROM ITS PRECURSOR BY EITHER TISSUE OR PLASMA
CC KALLIKREIN.
CC -I- MISCELLANEOUS: Rats express four types of kininogens: the
CC classical HMW and LMW kininogens produced by alternative splicing
CC of the same gene, and two additional LMW-like kininogens: T-I and
CC T-II.
CC -I- SIMILARITY: Contains 3 cystatin-like domains.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC EMBL; M11885; AAA41491.1; -.
DR PIR; B28055; B28055.
DR GlycoSuiteDB; P08932; -.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 3.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Glycoprotein; Plasma; Repeat; Vasodilator; Multigene family;
KW Thiol protease inhibitor; Bradykinin; Acute phase; Signal.
FT SIGNAL 1 18
FT CHAIN 19 430 KININOGEN, T-II.
FT CHAIN 19 375 HEAVY CHAIN.
FT PEPTIDE 376 386 T-KININ.
FT CHAIN 387 430 LIGHT CHAIN.
FT DOMAIN 19 135 CYSTATIN-LIKE 1.
FT DOMAIN 136 257 CYSTATIN-LIKE 2.
FT DOMAIN 258 375 CYSTATIN-LIKE 3.
FT DISULFID 28 404 INTERCHAIN (BY SIMILARITY).
FT DISULFID 83 94 BY SIMILARITY.
FT DISULFID 107 125 BY SIMILARITY.
FT DISULFID 141 144 BY SIMILARITY.
FT DISULFID 205 217 BY SIMILARITY.
FT DISULFID 228 247 BY SIMILARITY.
FT DISULFID 263 266 BY SIMILARITY.
FT DISULFID 327 339 BY SIMILARITY.
FT DISULFID 350 369 BY SIMILARITY.
FT CARBOHYD 82 82 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 126 126 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 326 326 N-LINKED (GLCNAC. .) (POTENTIAL).
SQ SEQUENCE 430 AA; 47524 MW; 43EDF02D1BF55076 CRC64;
Query Match 56.5%; Score 388; DB 1; Length 430;

```

Best Local Similarity 62.1%; Pred. No. 1.6e-29; Mismatches 29; Indels 0; Gaps 0;
Matches 72; Conservative 15; Mismatches 29; Indels 0; Gaps 0;

QY 3 GKPVQPTKICVCPDIPNSPELZETLTHITKLNAENATFPFKIDNVKARVQV 62
D 252 GDDUFLSLPKKFCPCPKNIPVDSPELKEALCHSIAQLNAGNHLFPFKIDTVKASQV 311
QY 63 AGKYFIDFVARETTCESKEEELTESCTKKGLQSLDCAEAVVVPWEKKIYPTV 118
D 312 AGTKVYEFIARETNCSTQNTLTDCTCKHGLQSLNCANVYMPREWNKVPTV 367

RESULT 9
KNT1 RAT
ID KNT1 RAT STANDARD; PRT; 430 AA.
AC P01048; P04081;
DT 01-NOV-1986 (Rel. 03, Created)
DT 01-NOV-1988 (Rel. 09, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE T-kininogen I precursor (Major acute phase protein) (Alpha-1-MAP)
DE (Thioistatin) [Contains: T-kinin].
GN MAPI.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OK NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=86008264; PubMed=2413018;
RA Furuto-Kato S., Matsumoto A., Kikamura N., Nakanishi S.;
RT "Primary structures of the MRNAs encoding the rat precursors for
RT bradykinin and T-kinin. Structural relationship of kininogens with
RT major acute phase protein and alpha 1-cysteine proteinase
RT inhibitor.";
RL J. Biol. Chem. 260:12054-12059(1985).
RN [2]
RP SEQUENCE OF 5-430 FROM N.A., AND PARTIAL SEQUENCE.
RX MEDLINE=86008266; PubMed=2413019;
RA Anderson K.P., Heath E.C.;
RT "The relationship between rat major acute phase protein and the
RT kininogens.";
RL J. Biol. Chem. 260:12065-12071(1985).
RN [3]
RP SEQUENCE OF 7-430 FROM N.A.
RX MEDLINE=85127561; PubMed=2578992;
RA Cole T., Inglis A.S., Roxburgh C.M., Howlett G.J., Schreiber G.;
RT "Major acute phase alpha 1-protein of the rat is homologous to bovine
RT kininogen and contains the sequence for bradykinin: its synthesis is
RT regulated at the mRNA level.";
RL FEBS Lett. 182:57-61(1985).
RN [4]
RP SEQUENCE OF 1-65 FROM N.A.
RX MEDLINE=87250580; PubMed=2439509;
RA Fung W.-P., Schreiber G.;
RT "Structure and expression of the genes for major acute phase alpha 1-
RT protein (thioistatin) and kininogen in the rat.";
RL J. Biol. Chem. 262:9298-9308(1987).
CC -1- FUNCTION: Kininogens are plasma glycoproteins with a number of
CC functions: (1) as precursor of the active peptide bradykinin they
CC effect smooth muscle contraction, induction of hypotension and
CC increase of vascular permeability. (2) they play a role in blood
CC coagulation by helping to position optimally prekallikrein and
CC factor XI next to factor XII. (3) They are inhibitor of thiol
CC proteases.
CC -1- SUBCELLULAR LOCATION: Extracellular.
CC -1- TISSUE SPECIFICITY: Plasma.
CC -1- INDUCTION: In response to an inflammatory stimulant. T-kininogen
CC II synthesis is induced and the plasma concentration of
CC T-kininogen I is raised.
CC -1- PTM: AS T-KININ IS PRECEDED BY A MET INSTEAD OF AN ARG OR LYS, IT
CC IS NOT RELEASED FROM ITS PRECURSOR BY EITHER TISSUE OR PLASMA
CC KALLIKREIN
CC -1- MISCELLANEOUS: Rats express four types of kininogens: the

CC classical HMW and LMW kininogens produced by alternative splicing
CC of the same gene, and two additional LMW-like kininogens: T-I and
CC T-II.
CC -1- SIMILARITY: Contains 3 cystatin-like domains.
CC -1- CAUTION: In addition to the conflicts described in the feature
CC table, Ref.2 sequence differs from that shown in positions 257,
CC 262, 268, 269, 295, 314, 315, 331, 332 and 389. In all those
CC positions the alternate amino acid is the one present in T-II
CC kininogen.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; M11883; AAA41489.1; -
CC EMBL; M11661; AAA41570.1; -
CC EMBL; M16454; AAA41568.1; -
CC EMBL; X02239; CAA326162.1; ALT_SEQ.
CC PIR; A01286; KGRTH.
CC PIR; A23897; A23897.
CC PIR; A27115; A27115.
CC GlycoSuiteDB; P01048; -
CC InterPro; IPR000010; Cystatin.
CC Pfam; PF00031; Cystatin; 3.
CC SMART; SM00043; CY_3.
CC PROSITE; PS00287; CYSTATIN; 2.
CC Glycoprotein; Plasma; Repeat; Vasodilator; Multigene family;
CC Thiol protease inhibitor; Bradykinin; Acute phase; Signal.
CC SIGNAL 1 18
CC CHAIN 19 430 KININOGEN, T-I.
CC CHAIN 19 375 HEAVY CHAIN.
CC PEPTIDE 376 386 T-KININ.
CC CHAIN 387 430 LIGHT CHAIN.
CC DOMAIN 19 135 CYSTATIN-LIKE 1.
CC DOMAIN 136 257 CYSTATIN-LIKE 2.
CC DOMAIN 258 375 CYSTATIN-LIKE 3.
CC DISULFID 28 404 INTERCHAIN (BY SIMILARITY).
CC DISULFID 83 94 BY SIMILARITY.
CC DISULFID 107 125 BY SIMILARITY.
CC DISULFID 141 144 BY SIMILARITY.
CC DISULFID 205 217 BY SIMILARITY.
CC DISULFID 228 247 BY SIMILARITY.
CC DISULFID 263 266 BY SIMILARITY.
CC DISULFID 327 339 BY SIMILARITY.
CC DISULFID 350 369 BY SIMILARITY.
CC CARBOHYD 82 82 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 126 126 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 168 168 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 204 204 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CARBOHYD 326 326 N-LINKED (GLCNAC. .) (POTENTIAL).
CC CONFLICT 26 28 LNC -> MDR (IN REF. 2).
CC CONFLICT 55 55 V -> L (IN REF. 2).
CC CONFLICT 61 61 E -> K (IN REF. 1).
CC CONFLICT 83 83 C -> Y (IN REF. 3).
CC CONFLICT 166 166 S -> F (IN REF. 2 AND 3).
CC CONFLICT 179 181 REV -> TKI (IN REF. 2).
CC CONFLICT 193 193 N -> D (IN REF. 2).
CC CONFLICT 212 212 S -> F (IN REF. 2).
CC CONFLICT 214 214 R -> H (IN REF. 3).
CC CONFLICT 229 229 T -> R (IN REF. 2).
CC CONFLICT 233 233 H -> Y (IN REF. 2).
CC CONFLICT 257 257 E -> S (IN REF. 2).
CC CONFLICT 262 262 N -> K (IN REF. 2).
CC CONFLICT 264 264 R -> F (IN REF. 2).
CC CONFLICT 268 268 RE -> KN (IN REF. 2).
CC CONFLICT 295 295 I -> L (IN REF. 2).
CC CONFLICT 314 315 VI -> TK (IN REF. 2).
CC CONFLICT 331 332 SK -> TN (IN REF. 2).
CC CONFLICT 389 389 R -> Q (IN REF. 2).

FT CONFLICT 414 414 R -> G (IN REF. 2 AND 3).
 FT CONFLICT 415 415 A -> L (IN REF. 2).
 FT CONFLICT 420 421 DH -> ER (IN REF. 3).
 FT CONFLICT 430 430 P -> S (IN REF. 1).
 SQ SEQUENCE 430 AA; 47715 MW; FAEBB78FAF4723C3 CRC64;
 Query Match 55.3%; Score 380; DB 1; Length 430;
 Best Local Similarity 62.1%; Pred. No. 9e-29;
 Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGPDRIPDTPNSPELEETLTHITKLAENNAATFYFKIDNVKARQVW 62
 DB 252 GDLFELLPKCKGPREIPVDSPELKEALGSHIAQNAOHNFYFKIDTVKKATSQV 311
 QY 63 AGKYFIDFVARETTCKESNEELTESCE---TKKLGSLDCNAEYVVPMEKITYPTV 118
 DB 312 AGYIVIEFIARETNSKSKETELTADCTKHLGSLNCNANVYMPENKVVPTV 367
 RESULT 10
 ID - CYTF_MOUSE STANDARD; PRT; 144 AA.
 AC 089098;
 DT 15-DEC-1998 (Rel. 37, Created)
 DT 15-DEC-1998 (Rel. 37, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Cystatin F precursor (Leukocystatin) (Cystatin 7) (Cystatin-like
 DE metastasis-associated protein) (CMAP).
 GN C57.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=98298157; PubMed=9632704;
 RA Halfon S., Ford J., Foster J., Dowling L., Lucian L., Sterling M.,
 RA Xu Y., Weiss M., Ikeda M., Liggett D., Helms A., Caux C., Lebecque S.,
 RA Hannum C., Menon S., McClanahan T., Gorman D., Zurawski G.;
 RT "Leukocystatin, a new class II cystatin expressed selectively by
 RT hematopoietic cells.";
 RL J. Biol. Chem. 273:16400-16408(1998).
 CC -!- FUNCTION: Inhibits papain and cathepsin L but with affinities
 CC lower than other cystatins. May play a role in immune regulation
 CC through inhibition of a unique target in the hematopoietic system.
 CC -!- SUBCELLULAR LOCATION: Secreted (Probable).
 CC -!- SIMILARITY: Belongs to the cystatin family.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.ebi.ac.uk/ebis/sequence/>
 CC or send an email to license@ebi.ac.uk).
 CC
 CC EMBL; AF031826; AAC40140.1; -;
 CC EMBL; AF031825; AAC40139.1; -;
 CC HSSP; P01034; 1G96.
 CC MGI; MGI:1298217; Cst7.
 CC InterPro; IPR000010; Cystatin.
 CC Pfam; PF00031; Cystatin; 1.
 CC SMART; SM00043; CY; 1.
 CC PROSITE; PS00287; CYSTATIN; FALSE NEG.
 KW Thiol protease inhibitor; Glycoprotein; Signal.
 FT SIGNAL 1 18
 FT CHAIN 19 144
 FT ACT_SITE 26 36
 FT SITE 80 84
 FT DISULFID 98 109
 FT DISULFID 123 143
 FT BY SIMILARITY.
 SQ SEQUENCE 144 AA; 16380 MW; B5837334C1B4A89C CRC64;

Query Match 25.0%; Score 171.5; DB 1; Length 144;
 Best Local Similarity 35.5%; Pred. No. 1.7e-09;
 Matches 39; Conservative 22; Mismatches 42; Indels 7; Gaps 3;
 QY 4 KDFVQPTKICVGPDRIPDTPNSPELEETLTHITKLAENNAATFYFKIDNVKARQVW 63
 DB 27 KDLI---SSVKGPGPKTIETNPGVLKAARHVSVEKFNCTNDIFLFKESHVSALVQVVK 83
 QY 64 GKKYFIDFVARETTCKESNEELTESCE---TKKLGSLDCNAEYVVPW 110
 DB 84 GLKTMLEVKIGRTTCRAKTHQL-DNCFOTNPALKRTLYCYSEVWVWP 132
 RESULT 11
 ID - CYTF_HUMAN STANDARD; PRT; 145 AA.
 AC 076096; Q9UED4;
 DT 15-DEC-1998 (Rel. 37, Created)
 DT 15-DEC-1998 (Rel. 37, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Cystatin F precursor (Leukocystatin) (Cystatin 7) (Cystatin-like
 DE metastasis-associated protein) (CMAP).
 GN C57.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=98406133; PubMed=9733783;
 RA Ni J., Fernandez M.A., Danielsson L., Chiklaku R.A., Zhang J.,
 RA Grubb A., Su J., Gentz R., Abrahamson M.;
 RT "Cystatin F is a glycosylated human low molecular weight cysteine
 RT proteinase inhibitor.";
 RL J. Biol. Chem. 273:24797-24804(1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=98298157; PubMed=9632704;
 RA Halfon S., Ford J., Foster J., Dowling L., Lucian L., Sterling M.,
 RA Xu Y., Weiss M., Ikeda M., Liggett D., Helms A., Caux C., Lebecque S.,
 RA Hannum C., Menon S., McClanahan T., Gorman D., Zurawski G.;
 RT "Leukocystatin, a new class II cystatin expressed selectively by
 RT hematopoietic cells.";
 RL J. Biol. Chem. 273:16400-16408(1998).
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Morita M., Arakawa H., Yoshiuchi N.;
 RT "Human homologue of murine CMAP.";
 RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20399571; PubMed=10945474;
 RA Morita M., Hara Y., Tamai Y., Arakawa H., Nishimura S.;
 RT "Genomic construct and mapping of the gene for CMAP
 RT (Leukocystatin/Cystatin F, CST7) and identification of a proximal
 RT novel gene, BSCV (C20orf3).";
 RL Genomics 67:87-91(2000).
 RN [5]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21638749; PubMed=11780052;
 RA Deloukas P., Matthews L.H., Ashurst J., Burton J., Gilbert J.G.R.,
 RA Jones M., Stavrides G., Almeida J.P., Babbage A.K., Baggeley C.L.,
 RA Bailey J., Barlow K.F., Bates K.N., Bead L.M., Beare D.M.,
 RA Beasley O.P., Bird C.P., Blakey S.E., Bridgman A.M., Brown A.J.,
 RA Buck D., Burrill W.D., Butler A.P., Carder C., Carter N.P., Clee C.M.,
 RA Clegg S., Cobley V.E., Collier R.E., Connor R.E., Corby N.R.,
 RA Coulson A., Coville G.J., Deadman R., Dhani P.D., Dunn M.,
 RA Ellington A.G., Frankland J.A., Fraser A., French L., Garner P.,
 RA Hammond D.V., Griffiths C., Griffiths M.N.D., Gwilliam R., Hall R.E.,
 RA Graham S., Harley J.L., Heath P.D., Ho S., Holden J.L., Howden P.J.,
 RA Huckle E., Hunt A.R., Hunt S.E., Jekosch K., Johnson C.M., Johnson D.,
 Kay M.P., Kimberley A.M., King A., Knights A., Laird G.K., Lawlor S.,

RA Levaeslao M.H., Leversha M.A., Lloyd C., Lloyd D.M., Lovell J.D.,
RA Marsh V.L., Martin S.L., McConachie L.J., McIay K., McMurray A.A.,
RA Milne S.A., Mistry D., Moore M.J.F., Mullikin J.C., Nickerson T.,
RA Oliver K., Parker A., Patel R., Pearce T.A.V., Peck A.I.,
RA Phillimore B.J.C.T., Prathalingam S.R., Plumb R.W., Ransay H.,
RA Rice C.M., Ross M.T., Scott C.E., Sehra H.K., Shownkeen R., Sims S.,
RA Skuce C.D., Smith M.L., Soderlund C., Steward C.A., Sulston J.E.,
RA Swann R.M., Sycamore A.C., Taylor R., Tee L., Thomas D.W., Thorpe A.,
RA Tracey A., Tromans A.C., Vaudin M., Wall M., Wallis J.M., Williams S.A.,
RA Whitehead S.L., Whittaker P., Willey D.L., Williams L., Williams S.A.,
RA Wilming L., Wray P.W., Hubbard T., Durbin R.M., Bentley D.R., Beck S.,
RA Rogers J.;
RA "The DNA sequence and comparative analysis of human chromosome 20.";
RL Nature 414:865-871(2001).
[6]
RP SEQUENCE FROM N.A.
RN TISSUE=Lung;
RC MEDLINE=22388257; PubMed=12477932;
RX Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci R.D., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S.J., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Yuzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Farney J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Souffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smal M.A.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -!- FUNCTION: Inhibits pepsin and cathepsin L but with affinities
CC lower than other cystatins. May play a role in immune regulation
CC through inhibition of a unique target in the hematopoietic system.
CC -!- SUBCELLULAR LOCATION: Secreted (Probable).
CC -!- TISSUE SPECIFICITY: Primarily expressed in peripheral blood cells
CC and spleen.
CC -!- SIMILARITY: Belongs to the cystatin family.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AF036342; AAC35747.1; --
CC EMBL; AF031824; AAC39788.1; --
CC EMBL; AB015225; BAA34941.1; ALT_INIT.
CC EMBL; AB029636; BAB11886.1; ALT_INIT.
CC EMBL; AL035661; CAB75498.1; --
CC EMBL; BC015507; AAB15507.1; ALT_INIT.
CC HSSP; P01034; IG96.
CC Genew; HGNC:2479; CST7.
CC MIM; 603253; --
CC GO; GO:0004869; F:cysteine protease inhibitor activity; TAS.
CC GO; GO:0006955; P:immune response; TAS.
CC InterPro; IPR000010; Cystatin.
CC Pfam; PF00031; cystatin; 1.
CC SMART; SM00043; Cy; 1.
CC PROSITE; PS00287; CYSTATIN; 1.
CC Thiol Protease inhibitor; Glycoprotein; Signal.
KW SIGNAL 1 19
FT CHAIN 20 145
FT ACT_SITE 37 37
FT REACTIVE_SITE

FT SITE 81 85
FT DISULFID 99 110
FT DISULFID 124 144
FT CARBOHYD 62 62
FT CARBOHYD 115 115
SQ SEQUENCE 145 AA; 16454 MW; B2CC4F76857CB0F CRC64;
Query Match 23.8%; Score 163.5; DB 1; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.9e-09;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;
QY 11 TKICVCGPDIPTNSPELEETLTHITKLNNATTFYFKIDNVKKARVQVVGKKYFID 70
DB 32 SRVKGFPTKINDPGVLQAARYSVKPNCTNDFLFXESRITRALVQIVKGLKYLE 91
QY 71 FVARETTCKESNEELTESCE---TKKLGQSLDCAEAVVYVPEKIKYPTVTVNWE 124
DB 92 VEIGRTTCKNQHLRL-DDCDFQTNHLTKQLTLCYSEVVVWPV-----LQHFE 138
RESULT 12
CYTC_MACMU STANDARD; PRT; 146 AA.
AC Q19092;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Cystatin C precursor.
GN CST3.
OS Macaca mulatta (Rhesus macaque).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutharia; Primates; Catarrhini; Cercopithecoidea;
OC Cercopithecoidea; Macaca.
OX NCBI_TaxID=9544;
RN [1]
RP MEDLINE=97054523; PubMed=898820;
RX Wei L.H., Walker L.C., Levy E.;
RA "Cystatin C. Icelandic-like mutation in an animal model of
RA cerebrovascular beta-amyloidosis.";
RL Stroke 27:2080-2085(1996).
CC -!- FUNCTION: As an inhibitor of cysteine proteinases, this protein is
CC thought to serve an important physiological role as a local
CC regulator of this enzyme activity.
CC -!- SIMILARITY: Belongs to the cystatin family.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; U51912; AAB64050.1; --
CC HSSP; P01034; IG96.
CC InterPro; IPR000010; Cystatin.
CC Pfam; PF00031; cystatin; 1.
CC SMART; SM00043; Cy; 1.
CC PROSITE; PS00287; CYSTATIN; 1.
CC Thiol protease inhibitor; Amyloid; Signal.
KW SIGNAL 1 26
FT CHAIN 27 146
FT ACT_SITE 37 37
FT SITE 81 85
FT DISULFID 99 109
FT DISULFID 123 143
SQ SEQUENCE 146 AA; 15857 MW; F0B3BB774A29DF26 CRC64;
Query Match 20.2%; Score 138.5; DB 1; Length 146;
Best Local Similarity 27.9%; Pred. No. 2.4e-06;
Matches 34; Conservative 25; Mismatches 52; Indels 11; Gaps 4;

QY 8 OPTKICVGPRLPTNSPLEETLTHITKLAENNAATFYKIDNVKKARQVQVAGKY 67
 Db 31 KPFR-LVGFMDASVEEGRALDFAVSEYNKASNDYHGRALQVVRKQIVAGVNY 88
 QY 68 FIDFVARETTCSKESNEELTESC---ETKLGQSLDCAEAVVWPWEKKIYPVTNVHWE 124
 Db 89 FLDELGRITCTK--TQPNLDNCFHEQHLKKAFCSPQIYVPHQ-----GTWLSKST 142
 QY 125 CE 126
 Db 143 CQ 144

RESULT 13
 CYTM_HUMAN STANDARD; PRT; 149 AA.
 AC Q15828;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Cystatin M precursor (Cystatin E).
 GN CST6.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=97150844; PubMed=8995380;
 RA Sotiropoulou G., Anisowicz A., Sager R.;
 RT "Identification, cloning, and characterization of cystatin M, a novel
 RT cysteine proteinase inhibitor, down-regulated in breast cancer.";
 RL J. Biol. Chem. 272:903-910(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=97256812; PubMed=9099741;
 RA Mi J., Abrahamson M., Zhang M., Fernandez M.A., Grubb A., Su J.,
 RA Yu G.L., Li Y., Parmelee D., Xing L., Coleman T.A., Gentz S.,
 RA Thotakura R., Nguyen N., Hesselberg M., Gentz R.;
 RT "Cystatin E is a novel human cysteine proteinase inhibitor with
 RT structural resemblance to family 2 cystatins.";
 RL J. Biol. Chem. 272:10853-10858(1997).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Prostate;
 RX MEDLINE=23388257; PubMed=12477932;
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Boudreau M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Huiy S.W.,
 RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahy J., Helton E., Kettner M., Madan A., Rodriguez S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grinchman J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalusz D.B.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [4]
 RP CHARACTERIZATION, AND TISSUE SPECIFICITY.
 RX MEDLINE=21246880; PubMed=11348457;
 RA Zeeuwen P.L., Van Vlijmen-Willems I.M., Jansen B.J., Sotiropoulou G.,
 RA Curfs J.H., Meis J.F., Janssen J.J., Van Ruissen F., Schalkwijk J.;
 RT "Cystatin M/E expression is restricted to differentiated epidermal
 RT keratinocytes and sweat glands: a new skin-specific proteinase

RT inhibitor that is a target for cross-linking by transglutaminase.";
 RL J. Invest. Dermatol. 116:693-701(2001).
 CC -!- FUNCTION: Shows moderate inhibition of cathepsin B but is not
 CC active against cathepsin C.
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- TISSUE SPECIFICITY: Restricted to the stratum granulosum of normal
 CC skin, the stratum granulosum/spinosum of psoriatic skin, and the
 CC secretory coils of eccrine sweat glands. Low expression levels are
 CC found in the nasal cavity.
 CC -!- PTM: Substrate for transglutaminases. Acts as an acyl acceptor but
 CC not as an acyl donor.
 CC -!- SIMILARITY: Belongs to the cystatin family.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC
 CC -----
 CC EMBL; U62800; RAB06566.1; -;
 CC EMBL; U81233; RAB61305.1; -;
 CC EMBL; BC011334; RAB61305.1; -;
 CC HSP; P01038; ICEW
 CC Genew; HGNC:2478; CST6.
 CC MIM; 601891; -;
 CC GO; GO:0004869; F: cysteine protease inhibitor activity; TAS.
 CC GO; GO:0007345; P: embryogenesis and morphogenesis; TAS.
 CC InterPro; IPR000010; Cystatin.
 CC Pfam; PF00031; cystatin; 1.
 CC SMART; SM00043; CY; 1.
 CC PROSITE; PS00287; CYSTATIN; 1.
 CC Thiol protease inhibitor; Signal; Glycoprotein.
 KW SIGNAL 1 28 PROBABLE.
 FT CHAIN 29 149 CYSTATIN M.
 FT ACT SITE 36 36 REACTIVE SITE.
 FT SITE 80 84 SECONDARY AREA OF CONTACT.
 FT DISULFID 98 113 BY SIMILARITY.
 FT DISULFID 126 146 BY SIMILARITY.
 FT CARBOHYD 137 137 N-LINKED (GLCNAC...) (POTENTIAL).
 SQ SEQUENCE 149 AA; 16511 MW; 2076A78BFC9FAC8C CRC64;
 Query Match 20.2%; Score 138.5; DB 1; Length 149;
 Best Local Similarity 31.5%; Pred. No. 2.4e-06;
 Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
 QY 10 PTKICVGPRLPTNSPLEETLTHITKLAENNAATFYKIDNVKKARQVQVAGKY 69
 Db 30 PQERMYGELRLDSPDPQVQKAAQAAVASYNMGNSIYYFRDTHIKAQSLVAGIKYFL 89
 QY 70 DFVARETTCSKE---SNEELTESCSTKLGQ--SIDCNAEVVVPWE 111
 Db 90 TWEMGSTDCRKTREVTGDHVDLT-TCLAAGAQQEKLRCDFEVLVVPWQ 136
 RESULT 14
 CYTC_BOVIN STANDARD; PRT; 148 AA.
 ID CYTC_BOVIN
 AC P01035;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 15-JUL-1999 (Rel. 38, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Cystatin C precursor (Colostrum thiol proteinase inhibitor).
 GN CST3.
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.; SEQUENCE OF 66-83, AND CHARACTERIZATION.
 RC TISSUE=Cerebrospinal fluid, and Choroid plexus;

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:06:08 ; Search time 13.716 Seconds
(without alignments)
890.662 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687
Sequence: 1 GSGKDFVQPTKICVGCPRD.....VPWEKKIYPTVTNNHWECEP 127

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 78:**

1: Pir1.*
2: Pir2.*
3: Pir3.*
4: Pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	618	90.0	427	1 KGHUL1	kininogen, LMW pre
2	618	90.0	644	1 KGHUL1	kininogen, HMW pre
3	440	64.0	436	1 KGBOL1	kininogen, LMW I p
4	440	64.0	621	1 KGBOL1	kininogen, HMW I p
5	413	60.1	434	1 KGBOL2	kininogen, LMW II
6	413	60.1	619	1 KGBOL2	kininogen, HMW II
7	410	59.7	433	2 A28055	K-kininogen, LMW I
8	410	59.7	639	2 A25486	kininogen, HMW I p
9	388	56.5	430	2 A23897	major acute phase
10	388	56.5	430	2 B28055	T-kininogen, LMW I
11	381	55.5	423	1 KGRTM	major acute phase
12	380	55.3	430	1 KGRTM	T-kininogen I prec
13	137.5	20.0	146	1 UDBO	cystatin - bovine
14	132.5	19.3	142	1 UDHU	cystatin C precurs
15	130	18.9	127	2 S07085	cystatin C precurs
16	129	18.8	120	2 S10587	cystatin C - rat
17	128	18.6	111	2 A28793	cystatin - puff ad
18	127.5	18.6	140	2 A36163	cystatin C precurs
19	124.5	18.1	141	2 B29632	cystatin SA precurs
20	118.5	17.2	139	1 UDCH	cystatin precurs
21	113	16.4	141	2 JQ1470	cystatin S precurs
22	112	16.3	91	2 S68034	T-kininogen (clone
23	112	16.3	91	2 S68035	T-kininogen (clone
24	111	16.2	111	1 JC2040	cystatin - chum sa
25	109.5	15.9	141	1 UDHP2	cystatin SN precurs
26	108.5	15.8	141	1 UDHP1	cystatin S precurs
27	107	15.6	139	2 T33740	hypothetical prote
28	106	15.4	132	2 JC4918	cystatin precurs
29	105.5	15.4	162	2 A43428	onchocystatin - ne

RESULT 1

KGHUL1

kininogen, LMW precursor [validated] - human

N;Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen

N;Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen

C;Species: Homo sapiens (man)

C;Date: 06-Jul-1982 #sequence revision 27-Nov-1985 #text change 08-Dec-2000

C;Accession: A01280; B25276; A27900; A27699; A31905; A34030

R;Ohkubo, I.; Kurachi, K.; Takasawa, T.; Shiohara, H.; Sasaki, M.

Biochemistry 23, 5691-5697, 1984

A;Title: Isolation of a human cDNA for alpha-2-thiol proteinase inhibitor and its identi

A;Reference number: A90490; MUID:85122621; PMID:6441591

A;Accession: A01280

A;Molecule type: mRNA

A;Residues: 1-427 <OHK>

A;Cross-references: GB:K02566; NID:g177889; PIDN:AAA35497.1; PID:g177890

R;Takagaki, Y.; Kitamura, N.; Nakanishi, S.

J. Biol. Chem. 260, 8601-8609, 1985

A;Title: Cloning and sequence analysis of cDNAs for human high molecular weight and low

A;Reference number: A92544; MUID:85234582; PMID:2989293

A;Accession: B25276

A;Molecule type: mRNA

A;Residues: 1-427 <TAK>

A;Cross-references: GB:M11437; NID:g186751; PIDN:AAB59551.1; PID:g386853

R;Lottspeich, F.; Kellermann, J.; Henschen, A.; Rauth, G.; Mueller-Esterl, W.

in Kinins IV, part A, Greenbaum, L.M., and Margolius, H.S., eds., pp.91-95, Plenum, New

A;Title: Amino acid sequence of the light chain of human low molecular mass kininogen.

A;Reference number: A27900

A;Accession: A27900

A;Molecule type: protein

A;Residues: 390-427 <LOT>

R;Mindrou, T.; Carretero, O.A.; Proud, D.; Walz, D.; Scicli, A.G.

Biochem. Biophys. Res. Commun. 152, 519-526, 1988

A;Title: A new kinin moiety in human plasma kininogens.

A;Reference number: A27699; MUID:88209021; PMID:3365237

A;Accession: A27699

A;Molecule type: protein

A;Residues: 380-389 <MIN>

R;Maeda, H.; Matsumura, Y.; Kato, H.

J. Biol. Chem. 263, 16051-16054, 1988

A;Title: Purification and identification of [hydroxypropyl(3)]-bradykinin in ascitic flu

A;Reference number: A31905; MUID:89034061; PMID:3182782

A;Accession: A31905

A;Molecule type: protein

A;Residues: 381-389 <MAE>

R;Sasaguri, M.; Ikeda, M.; Ideishi, M.; Arakawa, K.

Biochem. Biophys. Res. Commun. 150, 511-516, 1988

A;Title: Identification of [hydroxyproline(3)]-lysyl-bradykinin released from human plas

A;Reference number: A34030; MUID:88106632; PMID:33357729

A;Accession: A34030

A;Molecule type: protein

A;Residues: 380-389 <SAS>

A;Accession: A34030

A;Molecule type: protein

A;Residues: 380-389 <SAS>

A;Accession: A34030

A;Molecule type: protein

A;Residues: 380-389 <SAS>

R; Kitamura, N.; Kitagawa, H.; Fukushima, D.; Takagaki, Y.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 260, 8610-8617, 1985
A; Title: Structural organization of the human kininogen gene and a model for its evolution
A; Reference number: A92545; MUID: 85234583; PMID: 2989294
A; Contents: annotation; gene organization
R; Pierce, J. V., 1968
Fed. Proc. 27, 52-57, 1968
A; Title: Structural features of plasma kinins and kininogens.
A; Reference number: A91455; MUID: 90255622; PMID: 4952632
A; Contents: annotation; bradykinin
C; Comment: The LMW kininogen precursor is produced from the same gene as the HMW form (see A91455).
C; Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
C; Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, 1
xyproline residue is present in the kininogen prior to the release of bradykinin.
C; Genetics:
A; Gene: GDB:KNG
A; Cross-references: GDB:125256; OMIM:228960
A; Map position: 3q27-3q27
A; Introns: 65/3; 102/3; 131/1; 188/3; 224/3; 253/1; 310/3; 346/3; 375/3; 401/3
C; Superfamily: kininogen; cystatin homology
C; Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; glyco
F; 1-18/Domain: signal sequence #status predicted <SIG>
F; 19-427/Product: LMW prokininogen (kininogen I) #status predicted <MAT>
F; 19-389,390-427/Product: LMW kininogen II #status predicted <MAT2>
F; 19-379/Product: LMW kininogen heavy chain #status predicted <HCH>
F; 19-131/Domain: cystatin homology <CYL>
F; 142-253/Domain: cystatin homology <CY2>
F; 264-375/Domain: cystatin homology <CY3>
F; 380-389/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>
F; 381-389/Product: bradykinin (kallidin I) #status experimental <BDY>
F; 390-427/Product: LMW kininogen light chain #status experimental <LCH>
F; 19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status predicted
F; 28-407,83-94,107-126,142-145,206-218,229-248,264-267,328-340,351-370/Disulfide bonds:
F; 48,469,205,294/Binding site: carboxylate (Asn) (covalent) #status predicted
F; 379-380/Cleavage site: Met-Lys (kallikrein) #status experimental
F; 383/Modified site: 4-hydroxyproline (Pro) (partial) #status experimental
F; 389-390/Cleavage site: Arg-Ser (kallikrein) #status experimental
F; 401/Binding site: carboxylate (Thr) (covalent) #status absent

Query Match 90.0%; Score 618; DB 1; Length 427;
Best Local Similarity 100.0%; Pred. No. 3.6e-50;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTITITKLAENNATFYKIDNVKKARQVV 62
|||||
Db 253 GKDFVQPTKICVGCPRDIPNTSPLEETLTITITKLAENNATFYKIDNVKKARQVV 312
|||||

QY 63 AGKKYFIDFVARETTCSKESNEBELTSCETKKGQSLDCNDAEVYVPEKKIYPTV 118
|||||
Db 313 AGKKYFIDFVARETTCSKESNEBELTSCETKKGQSLDCNDAEVYVPEKKIYPTV 368
|||||

RESULT 2
KGHUI1
kininogen, HMW precursor [validated] - human
N; Alternate names: alpha-2-thiol proteinase inhibitor; prokininogen; prokininogen
N; Contains: bradykinin (kallidin I); HMW kininogen I; HMW kininogen II; low molecular we
C; Species: Homo sapiens (man)
C; Date: 28-May-1986 #sequence revision 28-May-1986 #text change 08-Dec-2000
C; Accession: A01279; A25276; S32422; A91153; A24871; A27899; A27699; A31905; A34030; S02
R; Okubo, I.; Kurachi, K.; Takasawa, T.; Shiohara, H.; Sasaki, M.
Biochemistry 23, 5651-5697, 1984
A; Title: Isolation of a human cDNA for alpha-2-thiol proteinase inhibitor and its ident
A; Reference number: A30490; MUID: 85122621; PMID: 6441591
A; Accession: A01279
A; Molecule type: mRNA
A; Residues: 1-389 <OHK>
A; Cross-references: GB:K02566; MID:g17789
R; Takagaki, Y.; Kitamura, N.; Nakanishi, S.
J. Biol. Chem. 260, 8601-8609, 1985
A; Title: Cloning and sequence analysis of cDNAs for human high molecular weight and low
A; Reference number: A92544; MUID: 85234582; PMID: 2989293
A; Accession: A25276

A; Molecule type: mRNA
A; Residues: 1-592, 'I', 594-644 <TAK>
A; Cross-references: GB:M11437; MID:g186751; PIDN:AAB59550.1; PID:g386852
R; Auerwald, E. A.; Roesler, D.; Mentele, R.; Assfalg-Wachleideit, I.
FEBS Lett. 321, 93-97, 1993
A; Title: Cloning, expression and characterization of human kininogen domain 3.
A; Reference number: S32422; MUID: 93223854; PMID: 8467916
A; Accession: S32422
A; Molecule type: mRNA
A; Residues: 'ANSM', 253-377 <AUS>
A; Note: differences are due to known cloning artifacts
R; Lottspeich, F.; Kellermann, J.; Henschel, A.; Foerbach, B.; Muller-Esterl, W.
Eur. J. Biochem. 152, 307-314, 1985
A; Title: The amino acid sequence of the light chain of human high-molecular-mass kininogen
A; Reference number: A91153; MUID: 86030270; PMID: 4054110
A; Accession: A91153
A; Molecule type: protein
A; Residues: 379-644 <LOT>
A; Note: the bradykinin sequence preceding the light chain sequence was not determined in
R; Kellermann, J.; Lottspeich, F.; Henschel, A.; Mueller-Esterl, W.
Eur. J. Biochem. 154, 471-478, 1986
A; Title: Completion of the primary structure of human high-molecular-mass kininogen. The
A; Reference number: A24871; MUID: 86108361; PMID: 3484703
A; Accession: A24871
A; Molecule type: protein
A; Residues: 'Z', 20-380 <KEU>
R; Kellermann, J.; Lottspeich, F.; Henschel, A.; Mueller-Esterl, W.
in Kinins IV, Greenbaum, L.M., and Margolius, H.S., ed., pp. 85-89, Plenum Press, New York
A; Title: Amino acid sequence of the light chain of human high molecular mass kininogen.
A; Reference number: A27899
A; Accession: A27899
A; Molecule type: protein
A; Residues: 379-389, K, 390-407, 'Q', 409-644 <KEL2>
R; Mindrou, T.; Carretero, O. A.; Proud, D.; Walz, D.; Scicli, A. G.
Biochem. Biophys. Res. Commun. 152, 519-526, 1988
A; Title: A new kinin moiety in human plasma kininogens.
A; Reference number: A27699; MUID: 88209021; PMID: 3365237
A; Accession: A27699
A; Molecule type: protein
A; Residues: 380-389 <MIN>
R; Maeda, H.; Matsumura, Y.; Kato, H.
J. Biol. Chem. 263, 16051-16054, 1988
A; Title: Purification and identification of [hydroxyprolyl(3)]bradykinin in ascitic fluid
A; Reference number: A31905; MUID: 89034061; PMID: 3182782
A; Accession: A31905
A; Molecule type: protein
A; Residues: 381-389 <MAB>
R; Sasaguri, M.; Ikeda, M.; Ideishi, M.; Arakawa, K.
Biochem. Biophys. Res. Commun. 150, 511-516, 1988
A; Title: Identification of [hydroxyproline(3)]-lysyl-bradykinin released from human plas
A; Reference number: A34030; MUID: 88106632; PMID: 3337729
A; Accession: A34030
A; Molecule type: protein
A; Residues: 380-389 <SAS>
R; Lenarcic, B.; Gabrijelcic, D.; Rozman, B.; Drobnic-Kosorok, M.; Turk, V.
Biochem. Biophys. Res. Commun. 150, 511-516, 1988
A; Title: Human Cope-Seyler 369, 257-261, 1988
A; Title: Human cathepsin B and cysteine proteinase inhibitors (CPIs) in inflammatory and
A; Reference number: S02482; MUID: 89076517; PMID: 3264507
A; Accession: S02482
A; Molecule type: protein
A; Residues: 1-19, 189-192; 310-314; 381-389 <LENI>
R; Kato, H.; Matsumura, Y.; Maeda, H.
FEBS Lett. 232, 252-254, 1988
A; Title: Isolation and identification of hydroxyproline analogues of bradykinin in human
A; Reference number: A61495; MUID: 88211869; PMID: 3366244
A; Accession: A61495
A; Molecule type: protein
A; Residues: 380-389 <KAT1>
A; Experimental source: urine
A; Note: this peptide had Pro-383 modified to 4-hydroxyproline
A; Accession: B61495
A; Molecule type: protein
A; Residues: 381-389 <KAT2>

A;Experimental source: urine
A;Note: this peptide had Pro-383 modified to 4-hydroxyproline
A;Accession: C61495
A;Molecule type: protein
A;Residues: 380-389 <KAT3>
R;Lenarcic, B.; Krasovec, M.; Ritonja, A.; Olafsson, I.; Turk, V.
FEBS Lett. 280, 211-215, 1991
A;Title: Inactivation of human cystatin C and kininogen by human cathepsin D.
A;Reference number: S14303; MUID:91192133; PMID:2013314
A;Accession: S14447
A;Molecule type: protein
A;Residues: 264-359, 'N', 361-375 <LEN2>
R;Little, S.S.; Johnson, D.A.
Biochem. J. 307, 341-346, 1995
A;Title: Human mast cell tryptase isoforms: separation and examination of substrate-specificity
A;Reference number: S55239; MUID:95251593; PMID:7733867
A;Accession: S55239
A;Molecule type: protein
A;Residues: 450-452, 'X', 454, 'X', 456 <LIT>
R;Strazek, J.; Maachi, F.; le Nguyen, D.; Becchi, M.; Heulin, M.H.; Nabet, P.; Belleville, J.
FEBS Lett. 373, 207-211, 1995
A;Title: Purification from human plasma of a tetrapeptide that potentiates insulin-like
A;Reference number: S68059; MUID:96033974; PMID:7589467
A;Accession: S68059
A;Molecule type: protein
A;Residues: 431-434 <STR>
R;Kitamura, N.; Kitagawa, H.; Fukushima, D.; Takagaki, Y.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 260, 8610-8617, 1985
A;Title: Structural organization of the human kininogen gene and a model for its evolution
A;Reference number: A92545; MUID:85234583; PMID:2989294
A;Contents: annotation; gene organization
R;Pierce, J.V.
Fed. Proc. 27, 52-57, 1968
A;Title: Structural features of plasma kinins and kininogens.
A;Reference number: A91455; MUID:90255622; PMID:4952632
A;Contents: annotation; bradykinin
C;Comment: The HMW kininogen precursor and the LMW form are produced from the same gene
C;Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
C;Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is im-
C;Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i-
C;Comment: The residue is present in the kininogen prior to the release of bradykinin.
A;Genetics:
A;Gene: GDB:XNG
A;Cross-references: GDB:125256; OMIM:228960
A;Map position: 3q27-3q27
A;Introns: 65/3; 102/3; 131/1; 188/3; 224/3; 253/1; 310/3; 346/3; 375/3
A;Superfamily: kininogen; cystatin homology
C;Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; dupli-
F;1-18/Domain: signal sequence #status experimental <SIG>
F;19-644/Product: HMW kininogen I (prokininogen) #status experimental <MAT1>
F;19-379,390-644/Product: HMW kininogen II #status experimental <MAT2>
F;19-379/Domain: HMW kininogen heavy chain #status experimental <HCH>
F;19-131/Domain: cystatin homology <CV1>
F;142-253/Domain: cystatin homology <CV2>
F;264-375/Domain: cystatin homology <CV3>
F;380-389/Product: lysyl-bradykinin (kallidin II) #status experimental <KB DY>
F;381-389/Product: bradykinin (kallidin I) #status experimental <BDY>
F;390-644/Domain: HMW kininogen light chain #status experimental <LCH>
F;421-510/Region: glycine/histidine/lysine-rich 30-residue repeats
F;431-434/Product: low molecular weight growth promoting factor #status experimental <GCH>
F;19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status experimental
F;28-614,83-94,107-126,142-145,206-218,229-248,264-267,328-340,351-370/Disulfide bonds:
F;48/Binding site: carbohydate (Asn) (covalent) #status absent
F;169,205,294/Binding site: carbohydate (Asn) (covalent) #status experimental
F;378-380/Cleavage site: Met-Lys (kallikrein) #status experimental
F;383/Modified site: 4-hydroxyproline (Pro) (partial) #status experimental
F;389-390/Cleavage site: Arg-Ser (kallikrein) #status experimental
F;408,533,542,546,557,571,593,628/Binding site: carbohydate (Thr) (covalent) #status ex-
F;577/Binding site: carbohydate (Ser) (covalent) #status experimental
Query Match 90.0%; Score 618; DB 1; Length 644;
Best Local Similarity 100.0%; Pred.No. 5.7e-50;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPPTKICVGCPDRDIPNTSPLESETLTHITTKLNENNAATFFPKIDNVKKARQVV 62
DB 253 GKDFVQPPTKICVGCPDRDIPNTSPLESETLTHITTKLNENNAATFFPKIDNVKKARQVV 312

QY 63 AGKYFIDFVARETTCSKESNEELTESCTKKLGQSLLDCNAAEVVVPWEKKIYPTV 118
DB 313 AGKYFIDFVARETTCSKESNEELTESCTKKLGQSLLDCNAAEVVVPWEKKIYPTV 368

RESULT 3
KGBOH1
kininogen, LMW I precursor - bovine
N;Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen
N;Contains: Bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C;Species: Bos primigenius taurus (cattle)
C;Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 22-Jun-1999
C;Accession: A01283
R;Kitamura, N.; Hirose, T.; Asai, M.; Inayama, S.; Nakanishi, S.
Proc. Natl. Acad. Sci. U.S.A. 80, 90-94, 1983
A;Title: Primary structures of bovine liver low molecular weight kininogen prepeptidase
A;Reference number: A93984; MUID:83117859; PMID:6572010
A;Accession: A01283
A;Molecule type: mRNA
A;Residues: 1-436 <NAB>
A;Cross-references: GB:J00010; GB:V00426; NID:g163256; PIDN:AAA30604.1; PID:g163256
C;Comment: The LMW kininogen precursor is produced from the same gene as the H₂O₂-stable bradykinin. Kininogen is a cysteine proteinase inhibitor, takes part in initiation
C;Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasoconstrictor. Bradykinin is present in the kininogen prior to the release of bradykinin.
C;Superfamily: kininogen; cystatin homology
C;Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor
F;1-18/Domain: signal sequence #status predicted <Sig>
F;19-436/Product: LMW kininogen I #status predicted <NAT>
F;19-378/Product: LMW kininogen I heavy chain #status predicted <HC>
F;19-130/Domain: cystatin homology <CY1>
F;141-252/Domain: cystatin homology <CV2>
F;263-374/Domain: cystatin homology <CV3>
F;379-388/Product: lysyl-bradykinin (kallidin II) #status predicted <KB DY>
F;380-388/Product: bradykinin (kallidin I) #status predicted <BD Y>
F;389-436/Product: LMW kininogen I light chain #status experimental <LCH>
F;19/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status unmodified
F;27-406 82-93, 106-125, 141-144, 205-217, 228-247, 263-266, 327-339, 350-369/Disulfide bonds (disulfide bonds) (covalent) #status predicted
F;47, 87, 168, 169, 197, 204/Binding site: carbonylhydrolase (Asn) (covalent) #status predicted
F;379/Cleavage site: Met-Lys (kallikrein) #status predicted
F;382/Modified site: 4-hydroxyproline (pro) #status predicted
F;388-389/Cleavage site: Arg-Ser (kallikrein) #status predicted

Query Match 64.0%; Score 440; DB 1; Length 436;
Best Local Similarity 70.4%; Pred. No. 1.7e-33;
Matches 81; Conservative 14; Mismatches 20; Indels 0; Gaps 0

QY 4 KDFVQPPTKICVGCPDRDIPNTSPLESETLTHITTKLNENNAATFFPKIDNVKKARQVWA 63
DB 253 KDFVQPPTKICVGCPDRDIPNTSPLESETLTHITTKLNENNAATFFPKIDNVKKARQVWA 312

QY 64 GKXYFIDFVARETTCSKESNEELTESCTKKLGQSLLDCNAAEVVVPWEKKIYPTV 118
DB 313 GLKYSIVFARETTCSKESNEELTKSCINHGQLHCDANVYVVPWEKKIYPTV 367

RESULT 4
KGBOH1
kininogen, HMW I precursor - bovine
N;Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen
N;Contains: Bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C;Species: Bos primigenius taurus (cattle)
C;Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 22-Jun-1999
C;Accession: A01281; A91923; A91938; A29559
R;Kitamura, N.; Takagaki, Y.; Furuto, S.; Tanaka, T.; Nawa, H.; Nakanishi, S.
Nature 305, 545-549, 1983
A;Title: A single gene for bovine high molecular weight and low molecular weight kininogen
A;Reference number: A93317; MUID:84014106; PMID:6571699

A:Accession: A01281
A:Molecule type: mRNA
A:Residues: 1-621 <KIT>
A:Cross-references: GB:V01491; GB:K01757; NID:G491; PIDN:CAA24735.1; PID:G492
R:Kato, H.; Nagasawa, S.; Suzuki, T.
J. Biochem. 67, 313-323, 1970
A:Title: Studies on the structure of bovine kininogen: cleavages of disulfide bonds and
A:Reference number: A91923; MUID:70180420; PMID:4986212
A:Accession: A91923
A:Molecule type: protein
A:Residues: 378-393 <KAT>
R:Han, Y.N.; Komiya, M.; Iwanaga, S.; Suzuki, T.
J. Biochem. 77, 55-68, 1975
A:Title: Studies on the primary structure of bovine high-molecular-weight kininogen. Ami
A:Reference number: A91938; MUID:75170265; PMID:1169237
A:Accession: A91938
A:Molecule type: protein
A:Residues: 458-498 <HAN>
R:Sueyoshi, T.; Miyata, T.; Hashimoto, N.; Kato, H.; Hayashida, H.; Miyata, T.; Iwanaga,
J. Biol. Chem. 262, 2768-2779, 1987
A:Title: Bovine high molecular weight kininogen. The amino acid sequence, positions of
A:Reference number: A92621; MUID:87137530; PMID:3546295
A:Accession: A29559
A:Molecule type: protein
A:Residues: 120-123, 125-127, 129-137 <SUE>
R:Kottspeich, F.; Kellermann, J.; Henschen, A.; Foertsch, B.; Muller-Esterl, W.
Eur. J. Biochem. 152, 307-314, 1985
A:Title: The amino acid sequence of the light chain of human high-molecular-mass kininogen
A:Reference number: A91153; MUID:86030270; PMID:4054110
A:Contents: annotation; bovine cleavage sites; bovine carbohydrate binding sites
R:Sueyoshi, T.; Miyata, T.; Kato, H.; Iwanaga, S.
Seikagaku 56, 808, 1984
A:Title: Disulfide bonds in bovine HMW kininogens.
A:Reference number: A94300
A:Contents: annotation; disulfide bonds
A:Note: article in Japanese
C:Comment: The HMW kininogen precursor is produced from the same gene as the LMW form as
C:Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
C:Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is im
C:Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i
xyproline residue is present in the kininogen prior to the release of bradykinin.
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; dupl
F:19-621/Product: HMW prokininogen I #status predicted <SIG>
F:19-379/Product: HMW kininogen I heavy chain #status experimental <HCH>
F:19-130/Domain: cystatin homology <CY1>
F:141-252/Domain: cystatin homology <CY2>
F:263-374/Domain: cystatin homology <CY3>
F:379-388/Product: bradykinin (kallidin II) #status experimental <KBDY>
F:388-621/Product: HMW kininogen I light chain #status experimental <BDY>
F:417-488/Region: glycine/histidine/lysine-rich
F:19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status experim
F:27-591, 82-93, 106-125, 141-144, 205-217, 228-247, 263-266, 327-339, 350-369/Disulfide bonds:
F:87, 168, 169, 204/Binding site: carbohydrate (Asn) (covalent) #status experimental
F:136/Binding site: carbohydrate (Thr) (covalent) (partial) #status experimental
F:197/Binding site: carbohydrate (Thr) (covalent) (partial) #status experimental
F:378-379/Cleavage site: Met-Lys (kallikrein) #status experimental
F:382/Modified site: 4-hydroxyproline (Pro) #status predicted
F:388-389/Cleavage site: Arg-Ser (kallikrein) #status experimental
F:398, 406, 512/Binding site: carbohydrate (Ser) (covalent) #status experimental
F:399, 400, 520, 524, 536, 548, 553, 570/Binding site: carbohydrate (Thr) (covalent) #status ex
F:498-499/Cleavage site: Arg-Thr (kallikrein) #status experimental

Query Match 64.0%; Score 440; DB 1; Length 621;
Best Local Similarity 70.4%; Pred. No. 2.4e-33;
Matches 81; Conservative 14; Mismatches 20; Indels 0; Gaps 0;

QY 4 KDFVOPPTKICVGCPRDPTNPSPELETLTITIKLNENNTATFYFKIDNVKKARQVVA 63
DB 253 KDFVOPPTLCAGCCKPIPVDSPLDEEPLSHSIKLNENHGDGAFYFKIDTVKKATVQVA 312

QY 64 GKYPFIDFVARETTCSKESNEELTESCETKKLGOSLDCNAEVYVVPWEKKIYPTV 118
DB 313 GLKYSIVFIARETTCSKSGSNEELTKSCEINIHGQILHCDANVYVVPWEKKIYPTV 367

RESULT 5
KGBOL2
kininogen, LMW II precursor - bovine
N:Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen
N:Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C:Species: Bos primigenius taurus (cattle)
C:Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 28-May-1999
C:Accession: A01284
R:Nawa, H.; Kitamura, N.; Hirose, T.; Asai, M.; Inayama, S.; Nakanishi, S.
Proc. Natl. Acad. Sci. U.S.A. 80, 90-94, 1983
A:Title: Primary structures of bovine liver low molecular weight kininogen precursors and
A:Reference number: A93984; MUID:83117859; PMID:6572010
A:Accession: A01284
A:Molecule type: mRNA
A:Residues: 1-434 <NAW>
A:Cross-references: GB:V00427; GB:J00011; NID:G489; PIDN:CAA23710.1; PID:G490
C:Comment: The LMW kininogen precursor is produced from the same gene as the HMW form as
C:Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
C:Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i
xyproline residue is present in the kininogen prior to the release of bradykinin.
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; glyco
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-434/Product: LMW kininogen II #status predicted <MAT>
F:19-377/Product: LMW kininogen I heavy chain #status predicted <HCH>
F:19-130/Domain: cystatin homology <CY1>
F:141-252/Domain: cystatin homology <CY2>
F:261-372/Domain: cystatin homology <CY3>
F:377-386/Product: lysyl-bradykinin (kallidin II) #status predicted <KBDY>
F:378-386/Product: bradykinin (kallidin I) #status predicted <BDY>
F:387-434/Product: LMW kininogen I light chain #status experimental <LCH>
F:13/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status predicted
F:27-404, 82-93, 106-125, 141-144, 205-217, 228-247, 261-264, 325-337, 348-367/Disulfide bonds:
F:47, 87, 168, 169, 197, 204, 280/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:376-377/Cleavage site: Met-Lys (kallikrein) #status predicted
F:380/Modified site: 4-hydroxyproline (Pro) #status predicted
F:386-387/Cleavage site: Arg-Ser (kallikrein) #status predicted

Query Match 60.1%; Score 413; DB 1; Length 434;
Best Local Similarity 67.2%; Pred. No. 5.5e-31;
Matches 78; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

QY 3 GKDFVOPPTKICVGCPRDPTNPSPELETLTITIKLNENNTATFYFKIDNVKKARQV 62
DB 252 GEDFL--PPMVCVGCCKPIPVDSPLDEEALNHSIAKLNENHGDGTFYFKIDTVKKATVQV 309

QY 63 AGKYPFIDFVARETTCSKESNEELTESCETKKLGOSLDCNAEVYVVPWEKKIYPTV 118
DB 310 GGLKYSIVFIARETTCSKSGSNEELTKSCEINIHGQILHCDANVYVVPWEKKIYPTV 365

RESULT 6
KGBOH2
kininogen, HMW II precursor - bovine
N:Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen
N:Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C:Species: Bos primigenius taurus (cattle)
C:Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 22-Jun-1999
C:Accession: A01282; A91923; A91941; A91938; B29559
R:Kitamura, N.; Takagaki, Y.; Furuto, S.; Tanaka, T.; Nawa, H.; Nakanishi, S.
Nature 305, 545-549, 1983
A:Title: A single gene for bovine high molecular weight and low molecular weight kininoge
A:Reference number: A93317; MUID:84014106; PMID:6571699
A:Accession: A01282
A:Molecule type: mRNA
A:Residues: 1-619 <KIT>
A:Cross-references: GB:V01492; GB:K01758; NID:G493; PIDN:CAA24736.1; PID:G494
R:Kato, H.; Nagasawa, S.; Suzuki, T.

R;Furuto-Kato, S.; Matsumoto, A.; Kitamura, N.; Nakanishi, S.
J. Biol. Chem. 260, 12054-12059, 1985
A;Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and
nibitor.
A;Reference number: A92496; MUID:86008264; PMID:2413018
A;Accession: A01426
A;Molecule type: mRNA
A;Residues: 1-430 <FUR>
A;Cross-references: GB:M11883; NID:G205084; PIDN:AAA41489.1; PID:G205085
R;Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 262, 2190-2198, 1987
A;Title: Differing expression patterns and evolution of the rat kininogen gene family.
A;Reference number: A92625; MUID:87137443; PMID:3029068
A;Accession: D25486
A;Molecule type: DNA
A;Residues: 375-430 <KIT>
R;Enjoji, K.; Kato, H.; Hayashi, I.; Oh-ishi, S.; Iwanaga, S.
J. Biol. Chem. 263, 973-979, 1988
A;Title: Purification and characterization of rat T-kininogens isolated from plasma of a
A;Reference number: A92729; MUID:88087226; PMID:3121623
A;Accession: A28526
A;Molecule type: protein
A;Residues: 'E', 20-48; 376-430 <ENJ>
R;Kanda, S.; Sugiyama, K.; Takahashi, M.; Shumiya, S.; Tomino, S.; Nagase, S.
Jpn. J. Cancer Res. 81, 63-68, 1990
A;Title: Identification of a protein increasing in serum of Nagase analbuminemic rats be
A;Reference number: P0193; MUID:90216390; PMID:2108948
A;Accession: P0193
A;Molecule type: mRNA
A;Residues: 330-420, 'R', 422-429, 'P' <KAN>
R;Anderson, K.P.; Croyle, M.L.; Lingrel, J.B.
Gene 81, 119-128, 1989
A;Title: Primary structure of a gene encoding rat T-kininogen.
A;Reference number: JQ0027; MUID:90034172; PMID:2806908
A;Accession: JQ0027
A;Molecule type: DNA
A;Residues: 1-60, 'B', 62-113, 'R', 115-165, 'F', 167-178, 'TKI', 182-211, 'P', 213-256, 'S', 258-38
R;Kageyama, R.; Kitamura, N.; Okubo, H.; Nakanishi, S.
J. Biol. Chem. 262, 2345-2351, 1987
A;Title: Differing utilization of homologous transcription initiation sites of rat K and
A;Reference number: A25488; MUID:87137465; PMID:3818598
A;Accession: B25488
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-48 <KAG>
A;Cross-references: GB:M14356; NID:G205090; PIDN:AAA41492.1; PID:G205091
R;Enjoji, K.; Kato, H.; Hayashi, I.; Oh-ishi, S.; Iwanaga, S.
J. Biol. Chem. 263, 965-972, 1988
A;Title: Purification and characterization of two kinds of low molecular weight kininoge
A;Reference number: A28525; MUID:88087225; PMID:3395530
A;Accession: A28525
A;Molecule type: protein
A;Residues: 376-430 <EN2>
R;Siererra, F.; Walter, R.; Vautravers, P.; Guigoz, Y.
Arch. Biochem. Biophys. 322, 333-338, 1995
A;Title: Identification of several isoforms of T-kininogen expressed in the liver of agi
A;Reference number: S68034; MUID:96032652; PMID:7574705
A;Accession: S68036
A;Molecule type: mRNA
A;Residues: 340-430 <SIE>
A;Experimental source: clone pSG17
C;Comment: At least three types of LMW kininogen precursors are present in rat plasma, t
ceding bradykinin.
C;Comment: T-kininogens contain T-kinin (I-S-bradykinin), a novel kinin isolated after t
d of an Arg or Lys, it is probably not released from its precursor by either tissue or p
C;Comment: The T-kininogens are produced in response to an inflammatory stimulant.
C;Genetics: 65/3, 102/3, 130/1, 187/3, 223/2, 252/1, 309/3, 345/3, 374/3, 398/3
A;introns: 65/3, 102/3, 130/1, 187/3, 223/2, 252/1, 309/3, 345/3, 374/3, 398/3
C;Superfamily: kininogen; cystatin homology
C;Keywords: acute phase; bradykinin; cysteine proteinase inhibitor; duplication; glycop
F;1-18/Domain: signal sequence; #status predicted <SIG>
F;19-430/Product: T-kininogen I #status experimental <NAT>

F;19-130/Domain: cystatin homology <CV1>
F;141-252/Domain: cystatin homology <CV2>
F;263-374/Domain: cystatin homology <CV3>
F;378-386/Product: bradykinin #status predicted <BDY>
F;19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status experiment
F;82,126,168,204,326/Binding site: carboxylate (Asn) (covalent) #status predicted
F;83-94,107-125,141-144,205-217,228-247,263-266,327-339,350-369/disulfide bonds: #status
Query Match 55.3%; Score 380; DB 1; Length 430;
Best Local Similarity 62.1%; Pred. No. 6,7e-28;
Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;
QY 3 GKDFVQPTKICVGCPRDIPNPSPELETLTHITKLAENNATFYFKIDNVKARQVW 62
DB 252 GDDLPELLPKNCRCGCPREIPVDSPELKEALGHSIAQLNAQNHIEYFKIDTVKKATSQV 311
QY 63 ACKKYFIDFVARETTCSEKNEELTESCTKLGSLDCNABVYVVPVEKKIYPTV 118
DB 312 AGVIVVIEFIARETNCOSKQSTELTADCTRHGLQSLNCNANVMRPENKVVPIV 367
RESULT 13
UDHU
N;Alternate names: thiol proteinase inhibitor
C;Species: Bos primigenius taurus (cattle)
C;Date: 28-Feb-1986 #sequence_revision 28-Feb-1986 #text_change 06-Dec-1996
C;Accession: A01271
R;Hirado, M.; Tsunawa, S.; Sakiyama, F.; Niinobe, M.; Fujii, S.
FEBS Lett. 186, 41-45, 1985
A;Title: Complete amino acid sequence of bovine colostrum low-M-r cysteine proteinase in
A;Reference number: A01271; MUID:85231205; PMID:3891407
A;Accession: A01271
A;Molecule type: protein
A;Residues: 1-112 <HIR>
C;Superfamily: cystatin; cystatin homology
C;Keywords: colostrum; cysteine proteinase inhibitor
F;2-112/Domain: cystatin homology <CVS>
F;48-52/Region: inhibitory #status predicted
F;66-76,90-110/disulfide bonds: #status predicted
Query Match 20.0%; Score 137.5; DB 1; Length 112;
Best Local Similarity 28.8%; Pred. No. 7.5e-06;
Matches 32; Conservative 25; Mismatches 35; Indels 19; Gaps 4;
QY 24 NSPELEETHTITKLAENNATFYFKIDNVKARQVQVAGKYFIDFVARETTCSEK 83
DB 12 NEEGVQEALSAFVSEFNEKSNDAVQSRVVRVVRARQVQVSGMNYFLDVLGRITCTK--S 69
QY 84 EELTESC-----ETKLGSLDCNABVYVVPVEKKIYPTVTVNWECE 126
DB 70 QANLDCPPHNQPHLREKL-----CSFQYVVPVWNV-----TINLVKFSQ 111
RESULT 14
UDHU
N;Alternate names: [validated] - human
C;Species: Homo sapiens (man)
C;Date: 06-Jul-1982 #sequence_revision 31-Mar-1991 #text_change 08-Dec-2000
C;Accession: S10216; S00004; J00095; A33400; S02751; A01270; A25434; S12288; A32732; A60
R;Abrahamson, M.; Olafsson, I.; Palsdottir, A.; Ulvbaeck, M.; Lundwall, A.; Jenson, O.
Biochem. J. 268, 287-294, 1990
A;Title: Structure and expression of the human cystatin C gene.
A;Reference number: S10216; MUID:90303202; PMID:2363674
A;Accession: S10216
A;Molecule type: DNA
A;Residues: 1-146 <AB1>
A;Cross-references: EMBL:X52255; NID:G30257; PIDN:CAA36497.1; PID:G296643
R;Abrahamson, M.; Grubb, A.; Olafsson, I.; Lundwall, A.
FEBS Lett. 216, 229-233, 1987
A;Title: Molecular cloning and sequence analysis of cDNA coding for the precursor of the
A;Reference number: S00004; MUID:87219149; PMID:3495457

A;Residues: 8-49 <ESN>
R;Esnard, A.; Esnard, F.; Guillou, F.; Gauthier, F.
FEBS Lett. 300, 131-135, 1992
A;Title: Production of the cysteine proteinase inhibitor cystatin C by rat Sertoli cells
A;Reference number: S21109; MUID:92225121; PMID:1563513
A;Accession: S21109
A;Molecule type: protein
A;Residues: 8, XX, 11-20 <ES2>
C;Superfamily: cystatin; cystatin homology
C;Keywords: cysteine proteinase inhibitor
F;16-127/Domain: cystatin homology <CYS>
F;80-90,104-124/Disulfide bonds: #status predicted

Query Match 18.9%; Score 130; DB 2; Length 127;
Best Local Similarity 28.0%; Pred. No. 4.3e-05;
Matches 30; Conservative 28; Mismatches 43; Indels 6; Gaps 4;
Qy 8 QPPTKICVGCPRDIPITNSPELEETLTHITITKLNANNATFYFKIDNVKKARVQVAGKKY 67
Db 11 RPPRL-LGAQEDADAGEGVQALDFAVSEYKNGSNDAYHSRAIQVVRARKQLVAGINY 69
Qy 68 PIDFVARETTTCKESNEELTESC---ETKKGSLDGNAEVYVYVPE 111
Db 70 YLDVEMGRITCTK-SQTNLT-NCPFHDPHLMRKALCSFQIYSVPWK 114

Search completed: September 24, 2004, 14:10:49
Job time : 14.716 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:08:41 ; Search time 44.704 Seconds
(without alignments)
913.519 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687
Sequence: 1 GSKGFQVPPTKICVGCPRD.....VPHEKKIYPTVTVNHWECEF 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1349238 seqs, 321558718 residues

Total number of hits satisfying chosen parameters: 1349238

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA:

1: /cgn2_6/ptodata/1/pubpaa/US07_PUBCOMB.pep.*
2: /cgn2_6/ptodata/1/pubpaa/PCT_NEW_PUB.pep.*
3: /cgn2_6/ptodata/1/pubpaa/US06_NEW_PUB.pep.*
4: /cgn2_6/ptodata/1/pubpaa/US06_PUBCOMB.pep.*
5: /cgn2_6/ptodata/1/pubpaa/US07_NEW_PUB.pep.*
6: /cgn2_6/ptodata/1/pubpaa/PCTUS_PUBCOMB.pep.*
7: /cgn2_6/ptodata/1/pubpaa/US08_NEW_PUB.pep.*
8: /cgn2_6/ptodata/1/pubpaa/US08_PUBCOMB.pep.*
9: /cgn2_6/ptodata/1/pubpaa/US09A_PUBCOMB.pep.*
10: /cgn2_6/ptodata/1/pubpaa/US09B_PUBCOMB.pep.*
11: /cgn2_6/ptodata/1/pubpaa/US09C_PUBCOMB.pep.*
12: /cgn2_6/ptodata/1/pubpaa/US09_NEW_PUB.pep.*
13: /cgn2_6/ptodata/1/pubpaa/US10A_PUBCOMB.pep.*
14: /cgn2_6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	618	90.0	390	15	US-10-162-335-82
2	618	90.0	398	15	US-10-162-335-70
3	618	90.0	427	10	US-09-949-039-29
4	618	90.0	615	15	US-10-162-335-72
5	618	90.0	644	15	US-10-162-335-74
6	618	90.0	644	15	US-10-162-335-84
7	381	55.5	424	14	US-10-316-253-217
8	380	55.3	430	14	US-10-316-253-215
9	163.5	23.8	145	14	US-10-329-428-2
10	163.5	23.8	167	10	US-09-746-783-197
11	163	23.7	178	9	US-09-969-834-1
12	138.5	20.2	121	9	US-09-775-932-14
13	138.5	20.2	128	9	US-09-775-932-12
14	138.5	20.2	149	9	US-09-940-497-2
15	137.5	20.0	112	8	US-08-849-303-16

16	137.5	20.0	112	16	US-10-655-136-16
17	136.5	19.9	118	9	US-09-775-932-24
18	132.5	19.3	120	9	US-09-775-932-2
19	132.5	19.3	120	16	US-10-695-194-2
20	132.5	19.3	146	8	US-08-849-303-17
21	132.5	19.3	146	9	US-09-940-497-3
22	132.5	19.3	146	9	US-09-969-834-3
23	132.5	19.3	146	14	US-10-329-428-3
24	132.5	19.3	146	14	US-10-376-564-47
25	132.5	19.3	146	16	US-10-655-136-17
26	132.5	19.3	146	16	US-10-695-194-1
27	132.5	19.3	249	16	US-10-257-384-A-2
28	132.5	19.3	641	16	US-10-257-384-A-4
29	131.5	19.1	317	12	US-10-210-172-82
30	131.5	19.1	345	12	US-10-210-172-86
31	131.5	19.1	356	12	US-10-210-172-84
32	131.5	19.1	369	12	US-10-210-172-78
33	131.5	19.1	369	12	US-10-210-172-80
34	131.5	19.1	382	12	US-10-315-664-93
35	131.5	19.1	382	12	US-09-978-360A-425
36	130	18.9	127	8	US-08-849-303-19
37	130	18.9	127	16	US-10-655-136-19
38	129.5	18.9	140	14	US-10-376-564-46
39	129.5	18.9	140	14	US-10-376-564-48
40	128	18.6	111	8	US-08-849-303-26
41	128	18.6	111	16	US-10-655-136-26
42	127.5	18.6	140	8	US-08-849-303-18
43	127.5	18.6	140	16	US-10-655-136-18
44	124.5	18.1	121	9	US-09-775-932-8
45	124.5	18.1	141	8	US-08-849-303-24

ALIGNMENTS

RESULT 1

US-10-162-335-82

; Sequence 82, Application US/10162335

; Publication No. US20040009480A1

; GENERAL INFORMATION:

; APPLICANT: Anderson, David W.

; APPLICANT: Baumgartner, Jason C.

; APPLICANT: Boldog, Ferenc L.

; APPLICANT: Casman, Stacie J.

; APPLICANT: Edinger, Shlomit R.

; APPLICANT: Gangolli, Esha A.

; APPLICANT: Gerlach, Valerie

; APPLICANT: Gorman, Linda

; APPLICANT: Guo, Xiaojia (Sasha)

; APPLICANT: Hjalte, Tord

; APPLICANT: Kekuda, Ramesh

; APPLICANT: Li, Li

; APPLICANT: MacDougall, John R.

; APPLICANT: Malyankar, Uriel M.

; APPLICANT: Miller, Isabelle

; APPLICANT: Padigaru, Muralidhara

; APPLICANT: Patturajan, Meera

; APPLICANT: Pena, Carol E. A.

; APPLICANT: Rastelli, Luca

; APPLICANT: Shimkets, Richard A.

; APPLICANT: Stone, David J.

; APPLICANT: Sytek, Kimberly A.

; APPLICANT: Vernet, Corine A. M.

; APPLICANT: Voss, Edward Z.

; APPLICANT: Zerhusen, Bryan D.

; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method

; FILE REFERENCE: 21402-377 B

; CURRENT APPLICATION NUMBER: US/10162,335

; PRIOR FILING DATE: 2002-10-01

; PRIOR APPLICATION NUMBER: 60/295,607

; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/295,661

; PRIOR FILING DATE: 2001-06-04

```

; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 82
; LENGTH: 390
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-82

Query Match          90.0%; Score 618; DB 15; Length 390;
Best Local Similarity 100.0%; Pred. No. 5.7e-59;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDVPQPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 62
Db 216 GKDVPQPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 275

QY 63 AGKGYFIDFVARETTCKSKESNEELTESCETKKGSLDCNAEYVVPWEKKIYPTV 118
Db 276 AGKGYFIDFVARETTCKSKESNEELTESCETKKGSLDCNAEYVVPWEKKIYPTV 331

RESULT 2
US-10-162-335-70
; Sequence 70, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Bolag, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Salomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjal, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malyankar, Uriel M.
; APPLICANT: Millet, Isabelle
; APPLICANT: Padigar, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shimkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; APPLICANT: Zernusen, Bryan D.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04

```

```

; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 70
; LENGTH: 398
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-70

Query Match          90.0%; Score 618; DB 15; Length 398;
Best Local Similarity 100.0%; Pred. No. 5.8e-59;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDVPQPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 62
Db 224 GKDVPQPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 283

QY 63 AGKGYFIDFVARETTCKSKESNEELTESCETKKGSLDCNAEYVVPWEKKIYPTV 118
Db 284 AGKGYFIDFVARETTCKSKESNEELTESCETKKGSLDCNAEYVVPWEKKIYPTV 339

RESULT 3
US-09-919-039-29
; Sequence 29, Application US/09919039
; Publication No. US20030108871A1
; GENERAL INFORMATION:
; APPLICANT: Kaser, Matthew R.
; TITLE OF INVENTION: GENES EXPRESSED IN TREATED HUMAN C3A LIVER CELL CULTURES
; FILE REFERENCE: PA-0035 US
; CURRENT APPLICATION NUMBER: US/09/919,039
; CURRENT FILING DATE: 2002-09-09
; PRIOR APPLICATION NUMBER: 60/222,113
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 401
; SOFTWARE: PERL Program
; SEQ ID NO 29
; LENGTH: 427
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. US20030108871A1 167507CD1
US-09-919-039-29

Query Match          90.0%; Score 618; DB 10; Length 427;
Best Local Similarity 100.0%; Pred. No. 6.4e-59;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDVPQPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 62
Db 253 GKDVPQPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 312

QY 63 AGKGYFIDFVARETTCKSKESNEELTESCETKKGSLDCNAEYVVPWEKKIYPTV 118
Db 313 AGKGYFIDFVARETTCKSKESNEELTESCETKKGSLDCNAEYVVPWEKKIYPTV 368

```


DB 284 AGKYPFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTV 339

RESULT 4
US-10-162-335-72
; Sequence 72, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalt, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malyankar, Uriel M.
; APPLICANT: Millet, Isabelle
; APPLICANT: Padigar, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shinkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; APPLICANT: Zerhusen, Bryan D.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 72
; LENGTH: 615
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-72

Query Match 90.0%; Score 618; DB 15; Length 615;
Best Local Similarity 100.0%; Pred. No. 1e-58;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GKDFVQPPKICVGCPRDIPNTSPLEELTHTITKLAENNATFYFKIDNVKKARVQV 62
DB 224 GKDFVQPPKICVGCPRDIPNTSPLEELTHTITKLAENNATFYFKIDNVKKARVQV 283
QY 63 AGKYPFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTV 118

DB 284 AGKYPFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTV 339

RESULT 5
US-10-162-335-74
; Sequence 74, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalt, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malyankar, Uriel M.
; APPLICANT: Millet, Isabelle
; APPLICANT: Padigar, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shinkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; APPLICANT: Zerhusen, Bryan D.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 74
; LENGTH: 644
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-74

Query Match 90.0%; Score 618; DB 15; Length 644;
Best Local Similarity 100.0%; Pred. No. 1.1e-58;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GKDFVQPPKICVGCPRDIPNTSPLEELTHTITKLAENNATFYFKIDNVKKARVQV 62
DB 253 GKDFVQPPKICVGCPRDIPNTSPLEELTHTITKLAENNATFYFKIDNVKKARVQV 312

QY 63 AGKYFIDFVARETTCSKESNEELTESCETKKGQSLDCNAEVVYVVPWEKKIYPTV 118
DB 313 AGKYFIDFVARETTCSKESNEELTESCETKKGQSLDCNAEVVYVVPWEKKIYPTV 368

RESULT 6

US-10-162-335-84

; Sequence 84, Application US/10162335

; Publication No. US20040009480A1

; GENERAL INFORMATION:

; APPLICANT: Anderson, David W.

; APPLICANT: Baumgartner, Jason C.

; APPLICANT: Boldog, Ferenc L.

; APPLICANT: Casman, Stacie J.

; APPLICANT: Edinger, Shlomit R.

; APPLICANT: Gangolli, Esha A.

; APPLICANT: Gerlach, Valerie

; APPLICANT: Gorman, Linda

; APPLICANT: Guo, Xiaojia (Sasha)

; APPLICANT: Hjal, Tord

; APPLICANT: Kekuda, Ramesh

; APPLICANT: Li, Li

; APPLICANT: MacDougall, John R.

; APPLICANT: Malyankar, Uriel M.

; APPLICANT: Millet, Isabelle

; APPLICANT: Padigaru, Muralidhara

; APPLICANT: Patturajan, Meera

; APPLICANT: Pena, Carol E. A.

; APPLICANT: Rastelli, Luca

; APPLICANT: Shimkets, Richard A.

; APPLICANT: Stone, David J.

; APPLICANT: Spytek, Kimberly A.

; APPLICANT: Vermet, Corine A. M.

; APPLICANT: Voss, Edward Z.

; APPLICANT: Zerhusen, Bryan D.

; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method

; CURRENT APPLICATION NUMBER: US/10/162,335

; CURRENT FILING DATE: 2002-10-01

; PRIOR APPLICATION NUMBER: 60/295,607

; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/295,661

; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/296,404

; PRIOR FILING DATE: 2001-06-06

; PRIOR APPLICATION NUMBER: 60/296,418

; PRIOR FILING DATE: 2001-06-06

; PRIOR APPLICATION NUMBER: 60/297,414

; PRIOR FILING DATE: 2001-06-11

; PRIOR APPLICATION NUMBER: 60/297,567

; PRIOR FILING DATE: 2001-06-12

; PRIOR APPLICATION NUMBER: 60/298,285

; PRIOR FILING DATE: 2001-06-14

; PRIOR APPLICATION NUMBER: 60/298,556

; PRIOR FILING DATE: 2001-05-15

; PRIOR APPLICATION NUMBER: 60/299,949

; PRIOR FILING DATE: 2001-06-21

; PRIOR APPLICATION NUMBER: 60/300,883

; PRIOR FILING DATE: 2001-08-26

; Remaining Prior Application data removed - See File Wrapper or PALM.

; NUMBER OF SEQ ID NOS: 201

; SEQ ID NO 84

; LENGTH: 644

; TYPE: PRT

; ORGANISM: Homo sapiens

US-10-162-335-84

Query Match

Best Local Similarity 90.0%; Score 618; DB 15; Length 644;

Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDIPNPSPELEETLTHITKLNANNATFYFKIDNVKARQV 62

DB 253 GKDFVQPTKICVGCPRDIPNPSPELEETLTHITKLNANNATFYFKIDNVKARQV 312
QY 63 AGKYFIDFVARETTCSKESNEELTESCETKKGQSLDCNAEVVYVVPWEKKIYPTV 118
DB 313 AGKYFIDFVARETTCSKESNEELTESCETKKGQSLDCNAEVVYVVPWEKKIYPTV 368

RESULT 7

US-10-316-253-217

; Sequence 217, Application US/10316253

; Publication No. US20030162706A1

; GENERAL INFORMATION:

; APPLICANT: The Procter & Gamble Company

; APPLICANT: Peters, Kevin

; APPLICANT: Thompson, Larry

; APPLICANT: Wang, Feng

; APPLICANT: Greis, Kenneth

; TITLE OF INVENTION: Angiogenesis Modulating Proteins

; FILE REFERENCE: 8865M

; CURRENT APPLICATION NUMBER: US/10/316,253

; CURRENT FILING DATE: 2002-12-10

; PRIOR APPLICATION NUMBER: US 60/355,295

; PRIOR FILING DATE: 2002-02-08

; NUMBER OF SEQ ID NOS: 308

; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 217

; LENGTH: 424

; TYPE: PRT

; ORGANISM: Rattus norvegicus

US-10-316-253-217

Query Match 55.5%; Score 381; DB 14; Length 424;

Best Local Similarity 62.1%; Pred. No. 5.8e-33;

Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDIPNPSPELEETLTHITKLNANNATFYFKIDNVKARQV 62

DB 246 GDDLPELLPKNCRGCPREIPVDSPELKEALGHISARLNAQHNIFFYFKIDTVKATQV 305

QY 63 AGKYFIDFVARETTCSKESNEELTESCETKKGQSLDCNAEVVYVVPWEKKIYPTV 118

DB 306 AGVIYVIERIARETNCSKQSKTELTADCTKHLGQSLCNCNANVYMPWENKVVPTV 361

RESULT 8

US-10-316-253-215

; Sequence 215, Application US/10316253

; Publication No. US20030162706A1

; GENERAL INFORMATION:

; APPLICANT: The Procter & Gamble Company

; APPLICANT: Peters, Kevin

; APPLICANT: Thompson, Larry

; APPLICANT: Wang, Feng

; APPLICANT: Greis, Kenneth

; TITLE OF INVENTION: Angiogenesis Modulating Proteins

; FILE REFERENCE: 8865M

; CURRENT APPLICATION NUMBER: US/10/316,253

; CURRENT FILING DATE: 2002-12-10

; PRIOR APPLICATION NUMBER: US 60/355,295

; PRIOR FILING DATE: 2002-02-08

; NUMBER OF SEQ ID NOS: 308

; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 215

; LENGTH: 430

; TYPE: PRT

; ORGANISM: Rattus norvegicus

US-10-316-253-215

Query Match

Best Local Similarity 55.3%; Score 380; DB 14; Length 430;

Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVCPDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKARQVV 62
Db 252 GDDIFELLPKNCRCFPIPDSPLEKALGHSAQLNAQHNIFFYFKIDTVKATQVV 311
QY 63 AGKYFIDFVARETTCKSKESNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTV 118
Db 312 AGVYVIEFIARETNCQSKSTELTADCTKHLGSLNCNANVTYRPNKRWPTV 367
RESULT 9
US-10-329-428-2
; Sequence 2, Application US/10329428
; Publication No. US20030114646A1
; GENERAL INFORMATION:
; APPLICANT: Li, et al.
; TITLE OF INVENTION: Human Cystatin F
; FILE REFERENCE: P2265P1D2
; CURRENT APPLICATION NUMBER: US/10/329,428
; PRIORITY FILING DATE: 2002-12-27
; PRIOR APPLICATION NUMBER: 60/014,795
; PRIOR FILING DATE: 1996-04-03
; PRIOR APPLICATION NUMBER: 08/832,535
; PRIOR FILING DATE: 1997-04-03
; PRIOR APPLICATION NUMBER: 09/019,485
; PRIOR FILING DATE: 1998-01-29
; PRIOR APPLICATION NUMBER: 09/528,436
; PRIOR FILING DATE: 2000-03-17
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: Patent in version 3.2
; SEQ ID NO 2
; LENGTH: 145
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-329-428-2
Query Match 23.8%; Score 163.5; DB 14; Length 145;
Best Local Similarity 31.6%; Pred No. 9.7e-10;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;
QY 11 TKICVCPDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKARQVVAGKYFID 70
Db 32 SRVKGPFKTIKNDPGVLQAARYSVKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 91
QY 71 FVARETTCKSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVVAHWE 124
Db 92 VEIGRTTCKKQHLRL-DDCDFQTNHTLKQTLSCYSEVWVWPV-----LQHEE 138
RESULT 10
US-09-746-783-197
; Sequence 197, Application US/09746783
; Publication No. US20030044935A1
; GENERAL INFORMATION:
; APPLICANT: Jacobs, Kenneth
; McCoy, John M.
; LaValle, Edward R.
; Racie, Lisa A.
; Treacy, Maurice
; Spaulding, Vikki
; Agostino, Michael J.
; Howes, Steven H.
; Fechtel, Kim
; TITLE OF INVENTION: SECRETED PROTEINS AND POLYNUCLEOTIDES
; ENCODING THEM
; NUMBER OF SEQUENCES: 231
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genetics Institute, Inc.
; STREET: 87 CambridgePark Drive
; CITY: Cambridge
; STATE: MA
; COUNTRY: U.S.A.
; ZIP: 02140
; COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA: US/09/746,783
FILING DATE: 21-Dec-2000
CLASSIFICATION: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Milasincic, Debra J.
REGISTRATION NUMBER: 46,931
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 227-7400
TELEFAX: (617) 742-4214
INFORMATION FOR SEQ ID NO: 197:
SEQUENCE CHARACTERISTICS:
LENGTH: 167 amino acids
TYPE: amino acid
STRANDEDNESS: <Unknown>
TOPOLOGY: linear
MOLECULE TYPE: protein
SEQUENCE DESCRIPTION: SEQ ID NO: 197:
US-09-746-783-197
Query Match 23.8%; Score 163.5; DB 10; Length 167;
Best Local Similarity 31.6%; Pred No. 1.2e-09;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;
QY 11 TKICVCPDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKARQVVAGKYFID 70
Db 54 SRVKGPFKTIKNDPGVLQAARYSVKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 113
QY 71 FVARETTCKSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVVAHWE 124
Db 114 VEIGRTTCKKQHLRL-DDCDFQTNHTLKQTLSCYSEVWVWPV-----LQHEE 160
RESULT 11
US-09-969-834-1
; Sequence 1, Application US/09969834
; Patent No. US20020102711A1
; GENERAL INFORMATION:
; APPLICANT: Bandman, Olga
; Goli, Surya K.
; TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE
; PROTEIN
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/969,834
FILING DATE: 01-Oct-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/09/471,765
FILING DATE: <Unknown>
APPLICATION NUMBER: US/08/791,522
FILING DATE: <Unknown>
APPLICATION NUMBER: 09/471,765
FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749

QY 3 GKDFVQPTKICVCPDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKARQVV 62
Db 252 GDDIFELLPKNCRCFPIPDSPLEKALGHSAQLNAQHNIFFYFKIDTVKATQVV 311
QY 63 AGKYFIDFVARETTCKSKESNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTV 118
Db 312 AGVYVIEFIARETNCQSKSTELTADCTKHLGSLNCNANVTYRPNKRWPTV 367
RESULT 9
US-10-329-428-2
; Sequence 2, Application US/10329428
; Publication No. US20030114646A1
; GENERAL INFORMATION:
; APPLICANT: Li, et al.
; TITLE OF INVENTION: Human Cystatin F
; FILE REFERENCE: P2265P1D2
; CURRENT APPLICATION NUMBER: US/10/329,428
; PRIORITY FILING DATE: 2002-12-27
; PRIOR APPLICATION NUMBER: 60/014,795
; PRIOR FILING DATE: 1996-04-03
; PRIOR APPLICATION NUMBER: 08/832,535
; PRIOR FILING DATE: 1997-04-03
; PRIOR APPLICATION NUMBER: 09/019,485
; PRIOR FILING DATE: 1998-01-29
; PRIOR APPLICATION NUMBER: 09/528,436
; PRIOR FILING DATE: 2000-03-17
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: Patent in version 3.2
; SEQ ID NO 2
; LENGTH: 145
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-329-428-2
Query Match 23.8%; Score 163.5; DB 14; Length 145;
Best Local Similarity 31.6%; Pred No. 9.7e-10;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;
QY 11 TKICVCPDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKARQVVAGKYFID 70
Db 32 SRVKGPFKTIKNDPGVLQAARYSVKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 91
QY 71 FVARETTCKSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVVAHWE 124
Db 92 VEIGRTTCKKQHLRL-DDCDFQTNHTLKQTLSCYSEVWVWPV-----LQHEE 138
RESULT 10
US-09-746-783-197
; Sequence 197, Application US/09746783
; Publication No. US20030044935A1
; GENERAL INFORMATION:
; APPLICANT: Jacobs, Kenneth
; McCoy, John M.
; LaValle, Edward R.
; Racie, Lisa A.
; Treacy, Maurice
; Spaulding, Vikki
; Agostino, Michael J.
; Howes, Steven H.
; Fechtel, Kim
; TITLE OF INVENTION: SECRETED PROTEINS AND POLYNUCLEOTIDES
; ENCODING THEM
; NUMBER OF SEQUENCES: 231
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genetics Institute, Inc.
; STREET: 87 CambridgePark Drive
; CITY: Cambridge
; STATE: MA
; COUNTRY: U.S.A.
; ZIP: 02140
; COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA: US/09/746,783
FILING DATE: 21-Dec-2000
CLASSIFICATION: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Milasincic, Debra J.
REGISTRATION NUMBER: 46,931
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 227-7400
TELEFAX: (617) 742-4214
INFORMATION FOR SEQ ID NO: 197:
SEQUENCE CHARACTERISTICS:
LENGTH: 167 amino acids
TYPE: amino acid
STRANDEDNESS: <Unknown>
TOPOLOGY: linear
MOLECULE TYPE: protein
SEQUENCE DESCRIPTION: SEQ ID NO: 197:
US-09-746-783-197
Query Match 23.8%; Score 163.5; DB 10; Length 167;
Best Local Similarity 31.6%; Pred No. 1.2e-09;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;
QY 11 TKICVCPDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKARQVVAGKYFID 70
Db 54 SRVKGPFKTIKNDPGVLQAARYSVKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 113
QY 71 FVARETTCKSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVVAHWE 124
Db 114 VEIGRTTCKKQHLRL-DDCDFQTNHTLKQTLSCYSEVWVWPV-----LQHEE 160
RESULT 11
US-09-969-834-1
; Sequence 1, Application US/09969834
; Patent No. US20020102711A1
; GENERAL INFORMATION:
; APPLICANT: Bandman, Olga
; Goli, Surya K.
; TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE
; PROTEIN
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/969,834
FILING DATE: 01-Oct-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/09/471,765
FILING DATE: <Unknown>
APPLICATION NUMBER: US/08/791,522
FILING DATE: <Unknown>
APPLICATION NUMBER: 09/471,765
FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749

```

; REFERENCE/DOCKET NUMBER: PP-0193 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 178 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; CLONE: 30443
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-969-834-1

Query Match      23.7%; Score 163; DB 9; Length 178;
Best Local Similarity 34.0%; Pred. No. 1.4e-09;
Matches 35; Conservative 20; Mismatches 44; Indels 4; Gaps 2;

QY 11 TKICVCGPRDIPNTPSPELEETLTHITIKLNAENNAFFYFKIDNVKARVQVVGKKYFI 70
Db 54 SRVPGFPKTIKNDGVIQAARISVEKENNCTNDFLESRTIRALVQIVKGLKYLE 113

QY 71 FVARETTCSKE---TKKLGQSLDCNAEVVVPWE 110
Db 114 VEIGRTTCKNQHLRL-DDCDFTQHTLAKTLSCYSEVVVVPW 155

RESULT 12
US-09-775-932-14
; Sequence 14, Application US/09775932
; Patent No. US20020137671A1
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent in Ver. 2.0
; SEQ ID NO 14
; LENGTH: 121
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-14

Query Match      20.2%; Score 138.5; DB 9; Length 121;
Best Local Similarity 31.5%; Pred. No. 4.2e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVCGPRDIPNTPSPELEETLTHITIKLNAENNAFFYFKIDNVKARVQVVGKKYFI 69
Db 2 PQERWVGELRDSPPDPQVQKAAQAAVSYNGSNSIYFRDTHIIKAQSQLVAGIKYFL 61

QY 70 DFVARETTCSKE---SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
Db 62 TMEMGSTDCRTRVTGDHVDLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 108

RESULT 13
US-09-775-932-12
; Sequence 12, Application US/09775932
; Patent No. US20020137671A1
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02

```

```

; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent in Ver. 2.0
; SEQ ID NO 12
; LENGTH: 128
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-12

Query Match      20.2%; Score 138.5; DB 9; Length 128;
Best Local Similarity 31.5%; Pred. No. 4.5e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVCGPRDIPNTPSPELEETLTHITIKLNAENNAFFYFKIDNVKARVQVVGKKYFI 69
Db 9 PQERWVGELRDSPPDPQVQKAAQAAVSYNGSNSIYFRDTHIIKAQSQLVAGIKYFL 68

QY 70 DFVARETTCSKE---SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
Db 69 TMEMGSTDCRTRVTGDHVDLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 115

RESULT 14
US-09-940-497-2
; Sequence 2, Application US/09940497
; Patent No. US20020052476A1
; GENERAL INFORMATION:
; APPLICANT: Ni et al.
; TITLE OF INVENTION: Human Cystatin E
; CURRENT APPLICATION NUMBER: US/09/940,497
; CURRENT FILING DATE: 2001-08-29
; PRIOR APPLICATION NUMBER: US 09/241,376
; PRIOR FILING DATE: 1999-02-02
; PRIOR APPLICATION NUMBER: US 08/744,138
; PRIOR FILING DATE: 1996-11-05
; PRIOR APPLICATION NUMBER: US 08/461,030
; PRIOR FILING DATE: 1995-06-05
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 2
; LENGTH: 149
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-940-497-2

Query Match      20.2%; Score 138.5; DB 9; Length 149;
Best Local Similarity 31.5%; Pred. No. 5.5e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVCGPRDIPNTPSPELEETLTHITIKLNAENNAFFYFKIDNVKARVQVVGKKYFI 69
Db 30 PQERWVGELRDSPPDPQVQKAAQAAVSYNGSNSIYFRDTHIIKAQSQLVAGIKYFL 89

QY 70 DFVARETTCSKE---SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
Db 90 TMEMGSTDCRTRVTGDHVDLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 136

RESULT 15
US-08-849-303-16
; Sequence 16, Application US/08849303
; Publication No. US20030221209A1
; GENERAL INFORMATION:
; APPLICANT: Atkinson, Howard J.
; APPLICANT: McPherson, Michael J.
; APPLICANT: Urwin, Peter E.
; TITLE OF INVENTION: MODIFIED PROTEINASE INHIBITORS
; NUMBER OF SEQUENCES: 79
; CORRESPONDENCE ADDRESS:

```

ADDRESSEE: Klauber & Jackson
STREET: 411 Hackensack Avenue, 4th Floor
CITY: Hackensack
STATE: New Jersey
COUNTRY: USA
ZIP: 07601
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/849,303
FILING DATE: 21-MAY-1997
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Jackson Esq., David A.
REGISTRATION NUMBER: 26,742
REFERENCE/DOCKET NUMBER: 1321-1-003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 201-487-5800
TELEFAX: 201-343-1684
TELEX: 133521
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 112 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
HYPOTHETICAL: NO
US-08-849-303-16

Query Match 20.0%; Score 137.5; DB 8; Length 112;
Best Local Similarity 28.8%; Pred. No. 4.9e-07;
Matches 32; Conservative 25; Mismatches 35; Indels 19; Gaps 4;
Qy 24 NSPELEETLTHTITKLNANNATFYFKIDNVKARVGVAGKGYFIDFVARETTCSKSN 83
Db 12 NEEGVQELSPAVSEFNKRSNDAYQSRVVRVRAKQVVGCMYFLDVELGRITCTK--S 69
Qy 84 BELTESC-----ETKKLQSLDCNAEVVVPWEKKIYPTVTVNHWECE 126
Db 70 QANLDCSPFHNPQLKREKL-----CSFYVVPWMN----TINLVKFSQC 111

Search completed: September 24, 2004, 14:13:04
Job time : 45.704 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

QM protein - protein search, using sw model

Run on: September 24, 2004, 14:07:01 ; Search time 14.268 Seconds
(without alignments)
445.051 Million cell updates/sec

Title: US-10-661-784-1
Perfect score: 660
Sequence: 1 GKDFVQPPKICVGCPRDIP.....YVPMEXKIYPTVNCQPLGM 123

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues
Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents AA.*
1: /cgn2_6/ptodata/2/iaa/5A COMB.pcp.*
2: /cgn2_6/ptodata/2/iaa/5B COMB.pcp.*
3: /cgn2_6/ptodata/2/iaa/6A COMB.pcp.*
4: /cgn2_6/ptodata/2/iaa/6B COMB.pcp.*
5: /cgn2_6/ptodata/2/iaa/PCTUS COMB.pcp.*
6: /cgn2_6/ptodata/2/iaa/backfiles1.pcp.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	600	90.9	117	1	US-08-193-114B-1
2	594	90.0	117	5	PCT-US92-06809-1
3	169	25.6	178	2	US-08-791-522-1
4	169	25.6	178	3	US-09-314-777-1
5	165.5	25.1	145	2	US-08-832-535-2
6	165.5	25.1	145	3	US-09-013-485-2
7	165.5	25.1	145	3	US-09-019-485-3
8	165.5	25.1	145	3	US-09-431-480-9
9	165.5	25.1	145	3	US-09-617-302-9
10	165.5	25.1	145	4	US-09-528-436B-2
11	155	23.5	27	3	US-08-676-242-11
12	145	22.0	64	3	US-08-676-242-22
13	145	22.0	64	3	US-09-402-732-1
14	138.5	21.0	121	4	US-09-775-932-14
15	138.5	21.0	128	4	US-09-775-932-12
16	138.5	21.0	149	2	US-08-461-030C-2
17	138.5	21.0	149	3	US-08-744-138-2
18	138.5	21.0	149	3	US-09-431-480-8
19	138.5	21.0	149	3	US-09-431-480-10
20	138.5	21.0	149	3	US-09-617-302-8
21	138.5	21.0	149	3	US-09-617-302-10
22	138.5	21.0	149	4	US-09-241-376-2
23	138.5	21.0	149	4	US-09-940-497-2
24	136	20.6	112	4	US-08-849-303-16
25	135	20.5	118	4	US-09-775-932-24
26	134	20.3	148	5	PCT-US95-07135-2
27	132	20.0	26	3	US-08-676-242-15

28	130	19.7	127	4	US-08-849-303-19	Sequence 19, Appl
29	129.5	19.6	140	4	US-08-866-319A-46	Sequence 46, Appl
30	129.5	19.6	140	4	US-08-866-319A-48	Sequence 48, Appl
31	129.5	19.6	382	4	US-09-589-360B-93	Sequence 93, Appl
32	128.5	19.5	146	6	5432264-6	Patent No. 5432264
33	128	19.4	111	4	US-08-849-303-26	Sequence 26, Appl
34	127.5	19.3	120	4	US-09-775-932-2	Sequence 2, Appl
35	127.5	19.3	140	3	US-09-431-480-5	Sequence 5, Appl
36	127.5	19.3	140	3	US-08-617-302-5	Sequence 5, Appl
37	127.5	19.3	140	4	US-08-849-303-18	Sequence 18, Appl
38	127.5	19.3	145	2	US-08-832-535-11	Sequence 11, Appl
39	127.5	19.3	146	2	US-08-791-522-3	Sequence 3, Appl
40	127.5	19.3	146	3	US-08-744-138-3	Sequence 3, Appl
41	127.5	19.3	146	3	US-09-019-485-4	Sequence 4, Appl
42	127.5	19.3	146	3	US-09-314-777-3	Sequence 3, Appl
43	127.5	19.3	146	3	US-09-431-480-6	Sequence 6, Appl
44	127.5	19.3	146	3	US-09-617-302-6	Sequence 6, Appl
45	127.5	19.3	146	4	US-09-241-376-3	Sequence 3, Appl

ALIGNMENTS

RESULT 1
US-08-193-114B-1
; Sequence 1, Application US/08193114B
; Patent No. 5472945
; GENERAL INFORMATION:
; APPLICANT: Schmaier, Alvin H.
; TITLE OF INVENTION: Modulation of Blood
; TITLE OF INVENTION: Pressure and Inhibition of Platelet Activation
; TITLE OF INVENTION: with Kininogen Fragment
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Seidel, Gonda, Lavorigna &
; ADDRESS: Monaco, P.C.
; STREET: 1800 Two Penn Center Plaza
; CITY: Philadelphia
; STATE: Pennsylvania
; COUNTRY: U.S.A.
; ZIP: 19102
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 720 Kb
; COMPUTER: IBM PS/2
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: WordPerfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/193,114B
; FILING DATE: 9 February 1994
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: U.S. Application
; APPLICATION NUMBER: Serial No. 5472945 07/744,545
; FILING DATE: 13 August 1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Monaco, Daniel A.
; REGISTRATION NUMBER: 30,480
; REFERENCE/DOCKET NUMBER: 6056-137 CII
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (215) 568-8383
; TELEFAX: (215) 568-5549
; TELEX: No. 5472345e
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 117 amino acids
; TYPE: peptide
; TOPOLOGY: linear

US-08-193-114B-1
Query Match 90.9%; Score 600; DB 1; Length 117;
Best Local Similarity 100.0%; Pred. No. 5.8e-59;
Matches 112; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 12 CVGCPDIPNPSPELEETLTHITIKLNAENNAFFYFKIDNVKARVQVWAGKYFIDFVA 71
 Db 1 CVGCPDIPNPSPELEETLTHITIKLNAENNAFFYFKIDNVKARVQVWAGKYFIDFVA 60
 QY 72 RETTCSKESNEBELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 123
 Db 61 RETTCSKESNEBELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 112

RESULT 2

PCT-US92-06809-1

; Sequence 1, Application PC/TUS9206809

; GENERAL INFORMATION:

; APPLICANT: Schmaier, Alvin H.

; APPLICANT: Jiang, Yongping

; TITLE OF INVENTION: Modulation of Blood

; TITLE OF INVENTION: Pressure by Altering Bradykinin Levels

; NUMBER OF SEQUENCES: 2

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Temple University - Of the

; ADDRESSEE: Commonwealth System of Higher Education

; STREET: 406 University Services

; CITY: Philadelphia

; STATE: Pennsylvania

; COUNTRY: U.S.A.

; ZIP: 19122

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Diskette, 3.50 inch, 720 Kb

; COMPUTER: IBM PS/2

; OPERATING SYSTEM: MS-DOS

; SOFTWARE: WordPerfect 5.1

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: PCT/US92/06809

; FILING DATE: 19910813

; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: U.S. Application

; APPLICATION NUMBER: Serial No. 744,545

; FILING DATE: 13 August 1991

; ATTORNEY/AGENT INFORMATION:

; NAME: Monaco, Daniel A.

; REGISTRATION NUMBER: 30,480

; REFERENCE/DOCKET NUMBER: 6056-137

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (215) 568-8383

; TELEFAX: (215) 568-5549

; TELEX:

; INFORMATION FOR SEQ ID NO: 1:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 117 amino acids

; TYPE: AMINO ACID

; TOPOLOGY: linear

PCT-US92-06809-1

Query Match

Best Local Similarity 90.0%; Score 594; DB 5; Length 117;

Matches 110; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 12 CVGCPDIPNPSPELEETLTHITIKLNAENNAFFYFKIDNVKARVQVWAGKYFIDFVA 71

Db 1 CVGCPDIPNPSPELEETLTHITIKLNAENNAFFYFKIDNVKARVQVWAGKYFIDFVA 60

QY 72 RETTCSKESNEBELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 123

Db 61 RETTCSKESNEBELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 112

RESULT 3

US-08-791-522-1

; Sequence 1, Application US/08791522

; Patent No. 5935817

; GENERAL INFORMATION:
 ; APPLICANT: Bandman, Olga
 ; APPLICANT: Goli, Surya K.
 ; TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE
 ; TITLE OF INVENTION: PROTEIN
 ; NUMBER OF SEQUENCES: 4
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Incyte Pharmaceuticals, Inc.
 ; STREET: 3174 Porter Drive
 ; CITY: Palo Alto
 ; STATE: CA
 ; COUNTRY: USA
 ; ZIP: 94304

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FastSeq for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/791,522

FILING DATE: Filed Herewith.

CLASSIFICATION: 514

PRIOR APPLICATION DATA:

APPLICATION NUMBER:

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Billings, Lucy J.

REGISTRATION/DOCKET NUMBER: 36,749

REFERENCE/DOCKET NUMBER: PF-0193 US

TELECOMMUNICATION INFORMATION:

TELEPHONE: 415-855-0555

TELEFAX: 415-845-4166

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 178 amino acids

TYPE: amino acid

STRANDEDNESS: single

TOPOLOGY: linear

IMMEDIATE SOURCE:

CLONE: 30443

US-08-791-522-1

Query Match

Best Local Similarity 25.6%; Score 169; DB 2; Length 178;

Matches 39; Conservative 22; Mismatches 49; Indels 10; Gaps 4;

QY 9 TKICVCPDIPNPSPELEETLTHITIKLNAENNAFFYFKIDNVKARVQVWAGKYFID 68

Db 54 SRVKGPFPTIKTNDPQVLAARYSVEKFNCTNDMLFKESRITRALVQIVKGLKYLE 113

QY 69 FVARETTCSKESNEBELTESCE---TKLGQSLDCNAEVVVPWEKKIYPTVN--CQPLGM 123

Db 114 VEIGRTTCKKNOHLEL-DDCDFQTNHTLKTLCSCYSEVWVVPW-----VPALRGACSPLSL 168

RESULT 4

US-09-314-777-1

; Sequence 1, Application US/09314777

; Patent No. 6110686

; GENERAL INFORMATION:

; APPLICANT: Bandman, Olga

; APPLICANT: Goli, Surya K.

; TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE

; TITLE OF INVENTION: PROTEIN

; NUMBER OF SEQUENCES: 4

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Incyte Pharmaceuticals, Inc.

; STREET: 3174 Porter Drive

; CITY: Palo Alto

; STATE: CA

; COUNTRY: USA

; ZIP: 94304

; COMPUTER READABLE FORM:


```

; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA: US/09/314,777
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/791,522
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0193 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 178 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; CLONE: 30443
;
US-09-314-777-1
Query Match 25.6%; Score 169; DB 3; Length 178;
Best Local Similarity 32.5%; Pred. No. 5.4e-11;
Matches 39; Conservative 22; Mismatches 49; Indels 10; Gaps 4;

QY 9 TKICVGCPRDPTNSPELEETLTHITIKLNAENNATFEKIDNVKARQVGVAGKYFID 68
DB 54 SRVKGPFPTIKTNDPGVLQAAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 113
QY 69 FVARETTCKESNEELTSCSCE---TKLIGQSLDCNAEVVVPWEKTIYPTVN--COPLGM 123
DB 114 VEIGRTTCKNQHLRL-DDCDFQTNHTLKQTLSCYSEVWVVPV---VPLRGAGCSPLSL 168

RESULT 5
US-08-832-535-2
; Sequence 2, Application US/08832535
; Patent No. 5919658
; GENERAL INFORMATION:
; APPLICANT: NI, JIAN
; APPLICANT: LI, HAODONG
; APPLICANT: YU, GUO-LIANG
; APPLICANT: GENTZ, REINER L
; TITLE OF INVENTION: HUMAN CYSTATIN F
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HUMAN GENOME SCIENCES, INC.
; STREET: 9410 KEY WEST AVENUE
; CITY: ROCKVILLE
; STATE: MD
; COUNTRY: US
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/832,535
; FILING DATE: 03-APR-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: KIMBALL, PAUL C.
; REGISTRATION NUMBER: 34,610
; REFERENCE/DOCKET NUMBER: PF265
; TELECOMMUNICATION INFORMATION:

QY 9 TKICVGCPRDPTNSPELEETLTHITIKLNAENNATFEKIDNVKARQVGVAGKYFID 68
DB 32 SRVKGPFPTIKTNDPGVLQAAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 91

us-10-661-784-1-rai
; TELEPHONE: (201) 994-1700
; TELEFAX: (201) 994-1744
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 145 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
;
US-08-832-535-2
Query Match 25.1%; Score 165.5; DB 2; Length 145;
Best Local Similarity 32.5%; Pred. No. 1e-10;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;

QY 9 TKICVGCPRDPTNSPELEETLTHITIKLNAENNATFEKIDNVKARQVGVAGKYFID 68
DB 32 SRVKGPFPTIKTNDPGVLQAAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 91
QY 69 FVARETTCKESNEELTSCSCE---TKLIGQSLDCNAEVVVPWEKTI-YPTVNC 118
DB 92 VEIGRTTCKNQHLRL-DDCDFQTNHTLKQTLSCYSEVWVVPVWLQHFVEFVLR 144

RESULT 6
US-09-019-485-2
; Sequence 2, Application US/09019485
; Patent No. 6066617
; GENERAL INFORMATION:
; APPLICANT: LI, HAODONG
; APPLICANT: YU, GUO-LIANG
; APPLICANT: GENTZ, REINER
; APPLICANT: NI, JIAN
; TITLE OF INVENTION: Cystatin F
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: MD
; COUNTRY: US
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/019,485
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Benson, Robert H.
; REGISTRATION NUMBER: 30,446
; REFERENCE/DOCKET NUMBER: PF265P1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 3013098504
; TELEFAX: 3013098439
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 145 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
;
US-09-019-485-2
Query Match 25.1%; Score 165.5; DB 3; Length 145;
Best Local Similarity 32.5%; Pred. No. 1e-10;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;

QY 9 TKICVGCPRDPTNSPELEETLTHITIKLNAENNATFEKIDNVKARQVGVAGKYFID 68
DB 32 SRVKGPFPTIKTNDPGVLQAAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 91
```


;; PRIOR FILING DATE: 1998-01-29
;; PRIOR APPLICATION NUMBER: 08/832,555
;; PRIOR FILING DATE: 1999-04-03
;; PRIOR APPLICATION NUMBER: 60/044,795
;; PRIOR FILING DATE: 1996-04-03
;; NUMBER OF SEQ ID NOS: 16
;; SOFTWARE: Patent in version 3.2
;; SEQ ID NO 2
;; LENGTH: 145
;; TYPE: PRT
;; ORGANISM: Homo sapiens
US-09-528-436B-2

Query Match 25.1%; Score 165.5; DB 4; Length 145;
Best Local Similarity 32.5%; Pred. No. 1e-10;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;
QY 9 TKICVGPDRPTNSPELEETLTITIKLNAENATYFKIDNVKKARVQVAGKYFID 68
DB 32 SRVXPGFPKTIKTNDPGVQAARYSVEKFNNTNDMELFKESRITRALVQIVKGLKYNLE 91
QY 69 FVARETTCKESNEELTESCS--TKKLQSLDCNAEVVVPWEKKI-YPTVNC 118
DB 92 VEIGRTCKCKQHLL-LDCCFQTNHILKQTLSCYSEVVVPVNLQHPFVPLRC 144

RESULT 11
US-08-676-242-11
; Sequence 11, Application US/08676242C
; Patent No. 6143719
; GENERAL INFORMATION:
; APPLICANT: The Regents of the University of Michigan
; APPLICANT: Schmaier, Alvin H.
; APPLICANT: Hasan, Ahmed A.K.
; TITLE OF INVENTION: Bradykinin Analogs As Selective Thrombin Inhibitors
; FILE REFERENCE: 8820-2 US
; CURRENT APPLICATION NUMBER: US/08/676,242C
; PRIOR FILING DATE: 2000-07-16
; EARLIER APPLICATION NUMBER: 60/000,096
; EARLIER FILING DATE: 1995-06-09
; PRIOR FILING DATE: 1996-06-07
; EARLIER FILING DATE: 1996-06-07
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: Patent in Ver. 2.1
; SEQ ID NO 11
; LENGTH: 27
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bradykinin
; OTHER INFORMATION: analog
US-08-676-242-11

Query Match 23.5%; Score 155; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 1.6e-10;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 97 LDCNAEVVVPWEKKIYPTVNCQPLGM 123
DB 1 LDCNAEVVVPWEKKIYPTVNCQPLGM 27

RESULT 12
US-08-676-242-22
; Sequence 22, Application US/08676242C
; Patent No. 6143719
; GENERAL INFORMATION:
; APPLICANT: The Regents of the University of Michigan
; APPLICANT: Schmaier, Alvin H.
; APPLICANT: Hasan, Ahmed A.K.
; TITLE OF INVENTION: Bradykinin Analogs As Selective Thrombin Inhibitors
; FILE REFERENCE: 8820-2 US
; CURRENT APPLICATION NUMBER: US/08/676,242C

;; CURRENT FILING DATE: 2000-07-16
;; EARLIER APPLICATION NUMBER: 60/000,096
;; EARLIER FILING DATE: 1995-06-09
;; EARLIER APPLICATION NUMBER: PCT/US96/03940
;; EARLIER FILING DATE: 1996-06-07
;; NUMBER OF SEQ ID NOS: 24
;; SOFTWARE: Patent in Ver. 2.1
;; SEQ ID NO 22
;; LENGTH: 64
;; TYPE: PRT
;; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: From human
; OTHER INFORMATION: kininogen heavy chain
US-08-676-242-22

Query Match 22.0%; Score 145; DB 3; Length 64;
Best Local Similarity 100.0%; Pred. No. 6.3e-09;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 99 CNAEVVVPWEKKIYPTVNCQPLGM 123
DB 1 CNAEVVVPWEKKIYPTVNCQPLGM 25

RESULT 13
US-09-402-732-1
; Sequence 1, Application US/09402732
; Patent No. 6251855
; GENERAL INFORMATION:
; APPLICANT: Schmaier, Alvin H.
; APPLICANT: Hasan, A.K. Ahmed
; TITLE OF INVENTION: Bradykinin Analogs As Selective Inhibitors of Cell
; TITLE OF INVENTION: Activation
; FILE REFERENCE: 8820-3
; CURRENT APPLICATION NUMBER: US/09/402,732
; CURRENT FILING DATE: 1999-12-01
; PRIOR APPLICATION NUMBER: 60/046,085
; PRIOR FILING DATE: 1997-04-23
; PRIOR APPLICATION NUMBER: PCT/US98/08015
; PRIOR FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: Patent in Ver. 2.1
; SEQ ID NO 1
; LENGTH: 64
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Segment of
; OTHER INFORMATION: human kininogen (residues 333-396 thereof)
US-09-402-732-1

Query Match 22.0%; Score 145; DB 3; Length 64;
Best Local Similarity 100.0%; Pred. No. 6.3e-09;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 99 CNAEVVVPWEKKIYPTVNCQPLGM 123
DB 1 CNAEVVVPWEKKIYPTVNCQPLGM 25

RESULT 14
US-09-775-932-14
; Sequence 14, Application US/09775932
; Patent No. 6534477
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:02:27 ; Search time 49.692 Seconds
(without alignments)
699.375 Million cell updates/sec

Title: US-10-661-784-1

Perfect score: 660

Sequence: 1 GKDFVQPTKICVGCPRDIP.....VVPWEKKIYPTVNCQLGM 123

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A_Geneseq_29Jan04:*

1: Geneseqp1980s:*

2: Geneseqp1990s:*

3: Geneseqp2000s:*

4: Geneseqp2001s:*

5: Geneseqp2002s:*

6: Geneseqp2003as:*

7: Geneseqp2003bs:*

8: Geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	660	100.0	123	3 AAY95426	Aay95426 Human hig
2	660	100.0	304	6 ABP70801	Abp70801 Human ext
3	660	100.0	322	6 ABP70799	Abp70799 Human ext
4	660	100.0	329	6 ABP92044	Abp92044 Human pro
5	660	100.0	358	6 ABP70800	Abp70800 Human ext
6	660	100.0	390	6 ABP99149	Abp99149 Novel hum
7	660	100.0	398	6 ABP99143	Abp99143 Novel hum
8	660	100.0	427	8 ADE76864	Adc76864 Human pro
9	660	100.0	615	6 ABP99144	Abp99144 Novel hum
10	660	100.0	626	5 ABP78707	Abp78707 Human hig
11	660	100.0	644	4 ABP21101	Abp21101 Novel hum
12	660	100.0	644	5 ABP78710	Abp78710 Human hig
13	660	100.0	644	6 ABP99150	Abp99150 Novel hum
14	660	100.0	644	6 ABP99145	Abp99145 Novel hum
15	628	95.2	122	3 AAB37447	Aab37447 Human kin
16	594	90.0	117	2 AAR33350	Aar33350 Domaine 3
17	589.5	89.3	435	4 ABG21105	Abg21105 Novel hum
18	477	72.3	436	1 AAP40257	Aap40257 Bradykini
19	453	68.6	357	6 ABA41202	Ab41202 Human DIT
20	450	68.2	434	1 AAP40633	Aap40633 Bradykini
21	409	62.0	235	5 ABG60077	Abg60077 Human DIT
22	320.5	48.6	248	4 ABG21102	Abg21102 Novel hum
23	316	47.9	369	4 ABG21099	Abg21099 Novel hum
24	190	28.8	305	4 ABG21100	Abg21100 Novel hum
25	175	26.5	167	2 AAW98907	Aaw98907 Mouse IMC

26	169	25.6	178	2 AAW69734	Aaw69734 Human cys
27	166	25.2	32	3 AAY95418	Aay95418 Anti-angi
28	165.5	25.1	126	3 AAB37445	Aab37445 Human cys
29	165.5	25.1	145	2 AAW32323	Aaw32323 Mature hu
30	165.5	25.1	145	2 AAW31502	Aaw31502 Human cys
31	165.5	25.1	145	2 AAY25708	Aay25708 Human cys
32	165.5	25.1	145	4 AAE02410	Aae02410 Human cys
33	165.5	25.1	145	4 AAE04439	Aae04439 Human cys
34	165.5	25.1	145	7 ADD14098	Add14098 Human src
35	165.5	25.1	167	2 AAY02287	Aay02287 Secreted
36	165.5	25.1	167	7 ADA45154	Ada45154 Human pol
37	161	24.4	32	3 AAY95408	Aay95408 Anti-angi
38	157	23.8	122	3 AAB37446	Aab37446 Human kin
39	156.5	23.7	167	2 AAW98910	Aaw98910 Mouse IMC
40	155	23.5	27	2 AAW54335	Aaw54335 Bradykini
41	153	23.2	27	3 AAY95425	Aay95425 Anti-angi
42	145	22.0	64	2 AAW54341	Aaw54341 Bradykini
43	145	22.0	64	2 AAW77418	Aaw77418 Kininogen
44	144.5	21.9	121	3 AAY81200	Aay81200 Human mut
45	144.5	21.9	128	3 AAY81189	Aay81189 Human mut

ALIGNMENTS

RESULT 1

AAY95426

ID AAY95426 standard; peptide; 123 AA.

XX AAY95426;

AC AAY95426;

XX AAY95426;

DT 25-SEP-2000 (first entry)

XX Human high mol.wt. kininogen domain 3.

DE Human high mol.wt. kininogen domain 3.

XX Anti-angiogenic; angiogenesis; inhibitor; kininogen; homologue;

KW endothelial cell proliferation; apoptosis; cancer; ocular disorder;

KW rheumatoid arthritis; cytostatic; antiarthritic; antirheumatic; therapy;

KW human; D3 peptide.

XX Homo sapiens.

OS Homo sapiens.

XX WO2000035407-A2.

PN 22-JUN-2000.

PD 22-DEC-1999; 99WO-US028465.

PF 16-DEC-1998; 98US-0112427P.

XX (UTEM) UNIV TEMPLE.

XX (MCCR/) MCCR R K.

XX Mcrae RK;

XX WPI; 2000-442247/38.

XX Composition for inhibiting angiogenesis and endothelial cell

PT proliferation, inducing endothelial cell apoptosis and treating cancer, 3

PT rheumatoid arthritis, and ocular disorders comprises a kininogen domain 3

XX Disclosure; Page 4; 44pp; English.

XX The present sequence is that of domain 3 of human high mol.wt. kininogen

CC (HK). The invention provides peptides (see AAY95405-24) that are

CC analogues of certain sites in the HK domain 3, specifically Asn275-

CC Lys282, Cys248-Cys249, Leu331-Tyr338 and Tyr299-Ser314. The peptides, in

CC which native Cys residues may be replaced by Ala residues, inhibit

CC endothelial cell proliferation and may also induce endothelial cell

CC apoptosis. Compositions including the peptides are used in claimed

CC methods for inhibiting angiogenesis, inhibiting endothelial cell

CC proliferation, and inducing endothelial cell apoptosis. Cancer.

CC rheumatoid arthritis, and ocular disorders characterized by undesired
 CC vascularization of the retina are treated
 XX
 SQ Sequence 123 AA;

Query Match 100.0%; Score 660; DB 3; Length 123;
 Best Local Similarity 100.0%; Pred. No. 2.3e-66;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNAFFYFKIDNVKKARQVQV 60
 DB 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNAFFYFKIDNVKKARQVQV 60
 QY 61 AGKXYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP 120
 DB 61 AGKXYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP 120
 QY 121 LGM 123
 DB 121 LGM 123

RESULT 2
 ABP70801
 ID ABP70801 standard; protein; 304 AA.
 XX
 AC ABP70801;
 XX
 DT 26-AUG-2003 (first entry)
 XX
 DE Human extracellular messenger, EXMES-28.
 XX
 KW Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO2003018612-A2.
 XX
 PD 06-MAR-2003.
 XX
 PF 22-AUG-2002; 2002WO-US027213.
 XX
 PR 24-AUG-2001; 2001US-0314811P.
 PR 14-DEC-2001; 2001US-0340584P.
 PR 18-JAN-2002; 2002US-0350595P.
 PR 11-MAR-2002; 2002US-0363432P.
 PR 15-MAR-2002; 2002US-0364607P.
 PR 05-APR-2002; 2002US-0370761P.
 PR 24-JUN-2002; 2002US-0391378P.

(INCY-) INCYTE GENOMICS INC.
 PA Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebajadian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PW;
 PI Ramkumar J;
 XX
 DR WPI; 2003-278643/27.
 DR N-PSDB; ACC42388.
 XX
 PT New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 PT disorders or cancer.
 XX
 PS Claim 1; Page 207; 224pp; English.
 XX
 CC The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to -28; ABP70774-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition

CC for diagnosing or treating a disease or condition associated with.
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer
 XX
 SQ Sequence 304 AA;

Query Match 100.0%; Score 660; DB 6; Length 304;
 Best Local Similarity 100.0%; Pred. No. 8e-66;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNAFFYFKIDNVKKARQVQV 60
 DB 130 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNAFFYFKIDNVKKARQVQV 189
 QY 61 AGKXYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP 120
 DB 190 AGKXYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP 249
 QY 121 LGM 123
 DB 250 LGM 252

RESULT 3
 ABP70799
 ID ABP70799 standard; protein; 322 AA.
 XX
 AC ABP70799;
 XX
 DT 26-AUG-2003 (first entry)
 XX
 DE Human extracellular messenger, EXMES-26.
 XX
 KW Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO2003018612-A2.
 XX
 PD 06-MAR-2003.
 XX
 PF 22-AUG-2002; 2002WO-US027213.
 XX
 PR 24-AUG-2001; 2001US-0314811P.
 PR 14-DEC-2001; 2001US-0340584P.
 PR 18-JAN-2002; 2002US-0350595P.
 PR 11-MAR-2002; 2002US-0363432P.
 PR 15-MAR-2002; 2002US-0364607P.
 PR 05-APR-2002; 2002US-0370761P.
 PR 24-JUN-2002; 2002US-0391378P.
 XX
 (INCY-) INCYTE GENOMICS INC.
 PA Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebajadian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PW;
 PI Ramkumar J;
 XX
 DR WPI; 2003-278643/27.
 DR N-PSDB; ACC42386.
 XX
 PT New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 PT disorders or cancer.
 XX
 PS Claim 1; Page 205-206; 224pp; English.
 XX
 CC The present invention relates to novel human extracellular messenger

CC proteins (EXMES-1 to-28; ABP70774-ABP70801) and their coding sequences
CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
CC for diagnosing or treating a disease or condition associated with
CC decreased expression or overexpression of functional EXMES e.g.
CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
CC cancer
XX
SQ Sequence 322 AA;
Query Match 100.0%; Score 660; DB 6; Length 322;
Best Local Similarity 100.0%; Pred. No. 8.7e-66; Indels 0; Gaps 0;
Matches 123; Conservative 0; Mismatches 0;
QY 1 GKDFVQPTKICVGCPRDIPTNSPELEETLTHITIKLNAENNATFYFKIDNVKARQVW 60
DB 148 GKDFVQPTKICVGCPRDIPTNSPELEETLTHITIKLNAENNATFYFKIDNVKARQVW 207
QY 61 AGKKYFIDFVARETTCSEKNEBELTESCTKLGSLDCNAEYVVPWEKKIYPTVNCOP 120
DB 208 AGKKYFIDFVARETTCSEKNEBELTESCTKLGSLDCNAEYVVPWEKKIYPTVNCOP 267
QY 121 LGM 123
DB 268 LGM 270
RESULT 4
ABU92044
ID ABU92044 standard; protein; 329 AA.
XX
AC ABU92044;
XX
DT 15-JUL-2003 (first entry)
XX
DE Human protein modification and maintenance molecule-24 (PMM-24).
XX
KW Human; protein modification and maintenance molecule; PMM; cancer;
KW cell proliferation disorder; atherosclerosis; neurological disorder;
KW epilepsy; Huntington's disease; stroke; immune disorder; allergy;
KW inflammatory disorder; AIDS; developmental disorder; hypothyroidism;
KW Cushing's syndrome; gastrointestinal disorder; epithelial disorder;
KW infection; cytostatic; antiarteriosclerotic; anticonvulsant; nootropic;
KW neuroprotective; cerebroprotective; anti-HIV; antiallergic; vulnary;
KW antiinflammatory; thyromimetic.
XX
OS Homo sapiens.
XX
XX WO2003031939-A2.
XX
XX 17-APR-2003.
XX
XX 11-OCT-2002; 2002WO-US032850.
XX
XX 12-OCT-2001; 2001US-0329689P.
XX
XX 25-OCT-2001; 2001US-0335703P.
XX
XX 09-NOV-2001; 2001US-0346887P.
XX
XX 28-NOV-2001; 2001US-0334145P.
XX
XX 06-DEC-2001; 2001US-0337451P.
XX
XX 14-DEC-2001; 2001US-0340584P.
XX
XX (INCY-) INCYTE GENOMICS INC.
XX
XX Ramkumar J, Gorvad AE, Baughn MR, Emerling BM, Yang J, Lee SY;
XX Tran UK, Becha SD, Duggan BM, Lee EA, Griffin JA, Li JX;
XX Sprague WW, Rafalia AJA, Chawla NK, Lehr-Mason PM, Kable AE, Yue H;
XX Marquis JP, Yao MG, Richardson TW, Tang TY, Jin P, Chien D;
XX Bhatia U, Burrill JD, Lee S, Blake JJ, Ho A, Zheng W,
XX WPI; 2003-430274/40.
XX
XX N-PSDB; ACA92439.
XX
XX New human protein modification and maintenance molecules (PMM), useful
XX for diagnosing, treating and preventing diseases or conditions associated

PT with the aberrant PMM expression e.g. cancer, atherosclerosis, or
PT infections.
XX
XX Claim 1; Page 264-265; 311pp; English.
XX
XX The present invention relates to the isolation of human protein
XX modification and maintenance molecules (PMM), and the polynucleotide
XX sequences encoding them. A total of 40 PMM polypeptides (designated PMM
XX -1 to PMM-40) are disclosed. The sequences of the invention are useful
XX for diagnosing a condition or disease associated with the expression of
XX PMM in a subject, preparing a polyclonal or monoclonal antibody, and
XX generating an expression profile of a sample containing the
XX polynucleotides. The diseases or conditions associated with decreased
XX expression or overexpression of PMM are cell proliferation disorders
XX (e.g. cancer, atherosclerosis), neurological disorders (e.g. epilepsy,
XX Huntington's disease, stroke), immune/inflammatory disorders, (e.g. AIDS,
XX allergies), developmental disorders (e.g. hypothyroidism, Cushing's
XX syndrome), gastrointestinal disorders (e.g. epithelial disorders, and infections. The
XX PMM polypeptides or their fragments are useful in screening compounds
XX for effectiveness as agonists or antagonists of the polypeptides, or in
XX altering the expression of the target polynucleotide and compounds that
XX specifically bind to, or modulate the activity of the polypeptide.
XX ABU92021-ABU92060 represent the human PMM polypeptides of the invention
XX
SQ Sequence 329 AA;
Query Match 100.0%; Score 660; DB 6; Length 329;
Best Local Similarity 100.0%; Pred. No. 9e-66;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GKDFVQPTKICVGCPRDIPTNSPELEETLTHITIKLNAENNATFYFKIDNVKARQVW 60
DB 155 GKDFVQPTKICVGCPRDIPTNSPELEETLTHITIKLNAENNATFYFKIDNVKARQVW 214
QY 61 AGKKYFIDFVARETTCSEKNEBELTESCTKLGSLDCNAEYVVPWEKKIYPTVNCOP 120
DB 215 AGKKYFIDFVARETTCSEKNEBELTESCTKLGSLDCNAEYVVPWEKKIYPTVNCOP 274
QY 121 LGM 123
DB 275 LGM 277
RESULT 5
ABP70800
ID ABP70800 standard; protein; 358 AA.
XX
XX AC ABP70800;
XX
XX 26-AUG-2003 (first entry)
XX
XX Human extracellular messenger, EXMES-27.
XX
XX Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
XX immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
XX endocrine disorder; cancer.
XX
XX OS Homo sapiens.
XX
XX WO2003018612-A2.
XX
XX 06-MAR-2003.
XX
XX 22-AUG-2002; 2002WO-US027213.
XX
XX 24-AUG-2001; 2001US-0314811P.
XX
XX 14-DEC-2001; 2001US-0340584P.
XX
XX 18-JAN-2002; 2002US-0350595P.
XX
XX 11-MAR-2002; 2002US-0363432P.
XX
XX 15-MAR-2002; 2002US-0364607P.
XX
XX 05-APR-2002; 2002US-0370761P.
XX
XX 24-JUN-2002; 2002US-0391378P.
XX

(INCYTE GENOMICS INC.)

PA Duggan BM, Lee S, Baughn MR, Hafalia AJA, Walia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebajadian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Ramkumar J;
 XX WPI: 2003-278643/27.
 DR N-PSDB; ACC42387.
 XX
 PT New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 PT disorders or cancer.
 XX
 PS Claim 1; Page 206; 224pp; English.
 XX
 CC The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to-28; ABP0774-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
 CC for diagnosing or treating a disease or condition associated with
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer
 XX
 SQ Sequence 358 AA;
 Query Match 100.0%; Score 660; DB 6; Length 358;
 Best Local Similarity 100.0%; Pred. No. 1e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFKIDNVKKARVQV 60
 DB 184 GKDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFKIDNVKKARVQV 243
 QY 61 AGKKYFIDFVARETTCKESNELTSCCTKLGSLDCNNAEVVVPVEKKIYPTVNCQP 120
 DB 244 AGKKYFIDFVARETTCKESNELTSCCTKLGSLDCNNAEVVVPVEKKIYPTVNCQP 303

QY 121 LGM 123
 DB 304 LGM 306

RESULT 6
 ABU99149
 ID ABU99149 standard; protein; 390 AA.
 AC ABU99149;
 XX
 DT 01-AUG-2003 (first entry)
 XX
 DE Novel human GPCR related protein NOV129.
 XX
 KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiant; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 XX
 CS Homo sapiens.
 XX
 XX WO200299116-A2.
 FN
 PD 12-DEC-2002.
 XX

PF 04-JUN-2002; 2002WO-US017428.
 XX
 PR 04-JUN-2001; 2001US-0295607P.
 PR 04-JUN-2001; 2001US-0295661P.
 PR 06-JUN-2001; 2001US-0296404P.
 PR 06-JUN-2001; 2001US-0296418P.
 PR 14-JUN-2001; 2001US-0298285P.
 PR 15-JUN-2001; 2001US-0298556P.
 PR 21-JUN-2001; 2001US-0299949P.
 PR 26-JUN-2001; 2001US-0300883P.
 PR 28-JUN-2001; 2001US-0301550P.
 PR 13-AUG-2001; 2001US-0311972P.
 PR 27-AUG-2001; 2001US-0315071P.
 PR 29-AUG-2001; 2001US-0315660P.
 PR 14-SEP-2001; 2001US-0322293P.
 PR 17-SEP-2001; 2001US-0322706P.
 PR 14-DEC-2001; 2001US-0341186P.
 PR 28-FEB-2002; 2002US-0361189P.
 PR 12-MAR-2002; 2002US-0363673P.
 PR 12-MAR-2002; 2002US-0363676P.
 PR 03-JUN-2002; 2002US-00363676.
 PA (CURA-) CURAGEN CORP.
 XX
 PI Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR;
 PI Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalte T, Kekuda R, Li L;
 PI Macdougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
 PI Pena CE, Rastelli L, Shinkets RA, Stone DU, Spytek KA, Vernet CAM;
 PI Voss EZ, Zerhusen BD;
 XX WPI: 2003-140627/13.
 DR N-PSDB; AC03653.
 XX
 PT New NOVX polypeptides and nucleic acids, useful for preventing or
 PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX
 PS Claim 1; Page 147; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95% identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX
 SQ Sequence 390 AA;
 Query Match 100.0%; Score 660; DB 6; Length 390;
 Best Local Similarity 100.0%; Pred. No. 1.1e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFKIDNVKKARVQV 60
 DB 216 GKDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFKIDNVKKARVQV 275

QY 61 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTVNCQP 120
 DB 276 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTVNCQP 335
 QY 121 LGM 123
 DB 336 LGM 338
 RESULT 7
 ABUS9143
 ID ABUS99143 standard; protein; 398 AA.
 AC ABUS99143;
 XX
 DT 01-AUG-2003 (first entry)
 XX
 DE Novel human GPCR related protein NOV12a.
 XX
 KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytostatic; cardiant; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOXV-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 XX
 OS Homo sapiens.
 XX
 WO200299116-A2.
 XX
 PD 12-DEC-2002.
 XX
 PF 04-JUN-2002; 2002WO-US017428.
 XX
 PR 04-JUN-2001; 2001US-0295607P.
 PR 04-JUN-2001; 2001US-0295661P.
 PR 06-JUN-2001; 2001US-0296404P.
 PR 06-JUN-2001; 2001US-0296418P.
 PR 14-JUN-2001; 2001US-0298285P.
 PR 15-JUN-2001; 2001US-0298558P.
 PR 21-JUN-2001; 2001US-0299949P.
 PR 26-JUN-2001; 2001US-0300883P.
 PR 28-JUN-2001; 2001US-0301550P.
 PR 13-AUG-2001; 2001US-0311972P.
 PR 27-AUG-2001; 2001US-0315071P.
 PR 29-AUG-2001; 2001US-0315660P.
 PR 14-SEP-2001; 2001US-0322293P.
 PR 17-SEP-2001; 2001US-0322708P.
 PR 14-DEC-2001; 2001US-0341186P.
 PR 28-FEB-2002; 2002US-0361189P.
 PR 12-MAR-2002; 2002US-0363673P.
 PR 12-MAR-2002; 2002US-0363676P.
 PR 03-JUN-2002; 2002US-00363676.
 XX
 PA (CURA-) CURAGEN CORP.
 XX
 PI Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR;
 PI Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalte T, Kekuda R, Li L;
 PI MacDougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
 PI Pena CE, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet CM;
 PI Voss EZ, Zerhusen BD;
 XX
 WIPI; 2003-140627/13.
 DR N-PSDB; ACD03647.
 DR
 XX
 PT New NOXV polypeptides and nucleic acids, useful for preventing or
 PT treating NOXV-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or

PT pharmacogenomics.
 XX
 PS Claim 1; Page 143; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOXV polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOXV-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOXV substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX
 SQ Sequence 398 AA;
 Query Match 100.0%; Score 660; DB 6; Length 398;
 Best Local Similarity 100.0%; Pred. No. 1.2e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPPKICVGCPRDIPNPSPELEETLTHITKLNANNATFYFKIDNVKKARVQVV 60
 DB 224 GKDFVQPPKICVGCPRDIPNPSPELEETLTHITKLNANNATFYFKIDNVKKARVQVV 283
 QY 61 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTVNCQP 120
 DB 284 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTVNCQP 343
 QY 121 LGM 123
 DB 344 LGM 346
 RESULT 8
 ADE76864
 ID ADE76864 standard; protein; 427 AA.
 XX
 AC ADE76864;
 XX
 DT 29-JAN-2004 (first entry)
 XX
 DE Human protein expressed in a liver disorder #9.
 XX
 KW human; liver disorder; hyperlipidaemia; hypertension; type II diabetes;
 KW tumour; liver; inflammatory disorder; immune response disorder;
 KW high-throughput screening; differential gene expression; gene therapy.
 XX
 OS Homo sapiens.
 XX
 FN US2003108871-A1.
 XX
 PD 12-JUN-2003.
 XX
 PF 30-JUL-2001; 2001US-00919039.
 XX
 XX 28-JUL-2000; 2000US-0222113P.
 XX
 PA (KASE/) KASER M R.
 XX
 PI Kaser MR;

XX WPI; 2004-031227/03.
 DR N-PSDB; ADE76863.
 XX
 PT Composition comprising several cDNAs that are differentially expressed in
 PT treated human C3A liver cell cultures, useful for treating liver
 PT disorders.
 XX
 PS Claim 1; SEQ ID NO 29; 41pp; English.
 XX
 CC The invention relates to a composition comprising several cDNAs that are
 CC differentially expressed in a liver disorder. The composition is useful
 CC for treating liver disorder such as hyperlipidaemia, hypertension, type
 CC II diabetes, tumours of the liver and disorders of the inflammatory and
 CC immune response. The composition is useful for a high-throughput method
 CC of screening several molecules or compounds to identify a ligand which
 CC specifically binds a cDNA. A protein encoded by the cDNA is useful for a
 CC high-throughput method for using a protein to screen several molecules or
 CC compounds to identify at least one ligand which specifically binds the
 CC protein which involves combining the protein encoded by the cDNA with
 CC several of molecules or compounds under conditions to allow specific
 CC binding, and detecting specific binding between the protein and a
 CC molecule or compound, therefore identifying a ligand which specifically
 CC binds the protein. The composition is useful for detecting and
 CC quantifying differential gene expression, can be used in gene therapy, to
 CC formulate prognosis and to design a treatment regimen and to monitor the
 CC efficacy of treatment. The present sequence represents the amino acid
 CC sequence of a protein encoded by a cDNA differentially expressed in a
 CC liver disorder.
 XX
 SQ Sequence 427 AA;
 Query Match 100.0%; Score 660; DB 8; Length 427;
 Best Local Similarity 100.0%; Pred. NO. 1.3e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPPKICVCGPRDIPNTSPLEETLTITIKLAENNAATFYFKIDNVKKARVQV 60
 DB 253 GKDFVQPPKICVCGPRDIPNTSPLEETLTITIKLAENNAATFYFKIDNVKKARVQV 312
 QY 61 AGKXIFDFVARETTCKSNBELTESCTKLGSLDCAAEVYVVPWEKKIYPTVNCQP 120
 DB 313 AGKXIFDFVARETTCKSNBELTESCTKLGSLDCAAEVYVVPWEKKIYPTVNCQP 372
 QY 121 LGM 123
 DB 373 LGM 375
 RESULT 9
 ID ABU99144
 AC ABU99144 standard; protein; 615 AA.
 XX
 AC ABU99144;
 XX
 DT 01-AUG-2003 (first entry)
 XX
 DE Novel human GPCR related protein NOV12b.
 KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiant; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 XX Homo sapiens.
 OS
 XX

PN W0200299116-A2.
 XX
 PD 12-DEC-2002.
 XX
 PF 04-JUN-2002; 2002WO-US017428.
 XX
 PR 04-JUN-2001; 2001US-0295607P.
 PR 04-JUN-2001; 2001US-0295661P.
 PR 06-JUN-2001; 2001US-0296404P.
 PR 06-JUN-2001; 2001US-0296418P.
 PR 14-JUN-2001; 2001US-0298285P.
 PR 15-JUN-2001; 2001US-0298556P.
 PR 21-JUN-2001; 2001US-0299949P.
 PR 26-JUN-2001; 2001US-0300893P.
 PR 28-JUN-2001; 2001US-0301550P.
 PR 13-AUG-2001; 2001US-0311972P.
 PR 27-AUG-2001; 2001US-0315071P.
 PR 29-AUG-2001; 2001US-0315660P.
 PR 14-SEP-2001; 2001US-0322293P.
 PR 17-SEP-2001; 2001US-0322706P.
 PR 14-DEC-2001; 2001US-0341186P.
 PR 28-FEB-2002; 2002US-0361189P.
 PR 12-MAR-2002; 2002US-0363673P.
 PR 12-MAR-2002; 2002US-0363676P.
 PR 03-JUN-2002; 2002US-00363676.
 XX (CURA-) CURAGEN CORP.
 PA
 XX Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR;
 PI Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalt T, Kekuda R, Li L;
 PI Macdougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
 PI Pena CE, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet CAM;
 PI Voss EZ, Zernhosen BD;
 XX WPI; 2003-140627/13.
 DR N-PSDB; ACD03648.
 XX
 XX New NOVX polypeptides and nucleic acids, useful for preventing or
 PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX
 PS Claim 1; Page 144; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX
 SQ Sequence 615 AA;
 Query Match 100.0%; Score 660; DB 6; Length 615;
 Best Local Similarity 100.0%; Pred. NO. 2.1e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVQPTKICVGCPRDIPNTSPSELEETLTHITITKLNNAENNAFYFKIDNVKKARVQV 60
 Db 224 GKDFVQPTKICVGCPRDIPNTSPSELEETLTHITITKLNNAENNAFYFKIDNVKKARVQV 283
 QY 61 AGKYFIDFVARETTCSKESNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTVNCOP 120
 Db 284 AGKYFIDFVARETTCSKESNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTVNCOP 343
 QY 121 LGM 123
 Db 344 LGM 346
 RESULT 10
 ABB78707
 ID ABB78707 standard; protein; 626 AA.
 XX
 AC ABB78707;
 XX
 DT 18-JUL-2002 (first entry)
 XX
 DE Human high molecular weight kininogen (HK) mature protein SEQ ID NO:1.
 XX
 KW Human; kininogen; high molecular weight kininogen; HK; D5 domain;
 KW D5 receptor; angiogenesis; endothelial cell; cytostatic; antitumour;
 KW antiatherosclerotic; vasotropic; vulnery; tranquilliser; thrombolytic;
 KW ophthalmological; gynaecological; antiulcer; antidiabetic; antiarthritic;
 KW antiangiogenic; antiapoptotic; endocrine; apoptosis; gene therapy.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT Domain 384..508
 FT /label= D5_domain
 FN WO200214369-A2.
 PD 21-FEB-2002;
 XX
 PF 24-JUL-2001; 2001WO-US023185.
 XX
 PR 24-JUL-2000; 2000US-0220194P.
 XX
 PA (ATTE-) ATTENUON LLC.
 XX
 PI Mazar AP, Juarez JC;
 XX
 DR WPI; 2002-393611/42.
 XX
 PT Novel human kininogen D5 domain polypeptides useful for treating
 PT conditions associated with endothelial cell migration, proliferation,
 PT invasion or angiogenesis, e.g. arthritis, macular degeneration, benign
 PT hyperplasia.
 XX
 PS Disclosure; Page 13; 84pp; English.
 XX
 CC The present invention describes an isolated polypeptide (I) that
 CC corresponds to the D5 domain of human kininogen, or biologically active
 CC peptide fragment, homologue or functional derivative, and which: (a)
 CC inhibits angiogenesis; (b) binds to the D5 binding site on endothelial
 CC cells (EC); (c) activates signalling pathways leading to the introduction
 CC of apoptosis in EC; and/or (d) inhibits the signalling pathway required
 CC for maintenance of EC viability. (I) has cytostatic, antitumour,
 CC antiatherosclerotic, vasotropic, vulnery, tranquilliser, thrombolytic,
 CC ophthalmological, gynaecological, antiulcer, antidiabetic, antiarthritic,
 CC antiangiogenic, antiapoptotic and endocrine activities. An antibody (IX)
 CC specific for an epitope of (I) is useful for inhibiting tumour growth or
 CC angiogenesis in a subject. (II), a D5 fusion polypeptide (II) or a dimeric
 CC or trimeric fusion polypeptide (III) can be used for inhibiting EC
 CC migration, proliferation, invasion, or angiogenesis, or for inducing EC
 CC apoptosis. An angiogenic EC-targeting pharmaceutical composition (X)
 CC comprising (I), (II), or (III), can be used for treating a subject having
 CC a disease or condition associated with undesired EC migration,

CC proliferation, invasion or angiogenesis. (I), (II), or (III) can be used
 CC for isolating a D5 domain binding molecule from a complex mixture and for
 CC isolating or enriching cells expressing D5 domain binding sites from a
 CC cell mixture. The present sequence represents the mature human high
 CC molecular weight kininogen (HK) protein, which is given in the
 CC exemplification of the present invention
 XX
 SQ Sequence 626 AA;

Query Match 100.0%; Score 660; DB 5; Length 626;
 Best Local Similarity 100.0%; Pred. No. 2.2e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVQPTKICVGCPRDIPNTSPSELEETLTHITITKLNNAENNAFYFKIDNVKKARVQV 60
 Db 235 GKDFVQPTKICVGCPRDIPNTSPSELEETLTHITITKLNNAENNAFYFKIDNVKKARVQV 294
 QY 61 AGKYFIDFVARETTCSKESNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTVNCOP 120
 Db 295 AGKYFIDFVARETTCSKESNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTVNCOP 354
 QY 121 LGM 123
 Db 355 LGM 357

RESULT 11

ID ABB21101 standard; protein; 644 AA.
 XX

AC ABB21101;
 XX

DT 18-FEB-2002 (first entry)
 XX

DE Novel human diagnostic protein #21092.
 XX

KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder.
 XX

OS Homo sapiens.
 XX

FN WO200175067-A2.
 XX

PD 11-OCT-2001.
 XX

PF 30-MAR-2001; 2001WO-US008631.
 XX

PR 31-MAR-2000; 2000US-00540217.
 XX

PR 23-AUG-2000; 2000US-00649167.
 XX

PA (HYSE-) HYSEQ INC.
 XX

PI Drmanac RT, Liu C, Tang YT;
 XX

DR WPI; 2001-639362/73.
 XX

DR N-PSDB; AAS85288.
 XX

PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX

PS Claim 20; SEQ ID NO 51460; 103pp; English.
 XX

CC The invention relates to isolated polynucleotide (I) and polypeptide (II)
 CC sequences. (I) is useful as hybridisation probes, polymerase chain
 CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
 CC and in recombinant production of (II). The polynucleotides are also used
 CC in diagnostics as expressed sequence tags for identifying expressed
 CC genes. (I) is useful in gene therapy techniques to restore normal
 CC activity of (II) or to treat disease states involving (II). (II) is
 CC useful for generating antibodies against it, detecting or quantitating a
 CC polypeptide in tissue, as molecular weight markers and as a food

CC supplement. (II) and its binding partners are useful in medical imaging
 CC of sites expressing (II). (I) and (II) are useful for treating disorders
 CC involving aberrant protein expression or biological activity. The
 CC polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. ABG00010-ABG30377 represent novel human diagnostic
 CC amino acid sequences of the invention. Note: The sequence data for this
 CC patent did not appear in the printed specification, but was obtained in
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 XX Sequence 644 AA;

Query Match 100.0%; Score 660; DB 4; Length 644;
 Best Local Similarity 100.0%; Pred. No. 2.3e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKARVQV 60
 Db 253 GKDFVQPPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKARVQV 312
 QY 61 AGKKYFDVFARETTCKESNEELTESCTKLGQSLDCNAEYVVPWEKKIYPTVNCQ 120
 Db 313 AGKKYFDVFARETTCKESNEELTESCTKLGQSLDCNAEYVVPWEKKIYPTVNCQ 372
 QY 121 LGM 123
 Db 373 LGM 375

RESULT 12

ABB78710
 ID ABB78710 standard; protein; 644 AA.

AC ABB78710;

XX 18-JUL-2002 (first entry)

DE Human high molecular weight kininogen (HK) protein.

DE Human; kininogen; high molecular weight kininogen; HK; D5 domain;
 KW D5 receptor; angiogenesis; endothelial cell; cytostatic; antitumour;
 KW antiatherosclerotic; vasotropic; vulnary; tranquilliser; thrombolytic;
 KW ophthalmological; gynaecological; antiulcer; antidiabetic; antiarthritic;
 KW antiangiogenic; antiapoptotic; endocrine; apoptosis; gene therapy.

OS Homo sapiens.

XX Key Location/Qualifiers

FT Peptide 1..18

FT Protein 19..644

FT /label= signal

FT /label= mature_human_high_molecular_weight_kininogen

FT Disulfide-bond 28..614

FT Disulfide-bond 83..94

FT Disulfide-bond 107..126

FT Disulfide-bond 142..145

FT Disulfide-bond 206..218

FT Disulfide-bond 229..248

FT Disulfide-bond 264..267

FT Disulfide-bond 328..340

FT Disulfide-bond 351..370

FT Domain 402..526

FT /label= D5_domain

XX WO200214369-A2.

XX 21-FEB-2002.

XX 24-JUL-2001; 2001WO-US023185.

PR 24-JUL-2000; 2000US-0220194P.

PA (ATTE-) ATTENUON LLC.

PI Mazar AP, Juarez JC;

XX WPI; 2002-393611/42.

XX Novel human kininogen D5 domain polypeptides useful for treating

PT conditions associated with endothelial cell migration, proliferation,

PT invasion or angiogenesis, e.g. arthritis, macular degeneration, benign

PT hyperplasia.

XX Disclosure; Fig 1B-E; 84pp; English.

XX The present invention describes an isolated polypeptide (I) that

CC corresponds to the D5 domain of human kininogen, or biologically active

CC peptide fragment, homologue or functional derivative, and which: (a)

CC inhibits angiogenesis; (b) binds to the D5 binding site on endothelial

CC cells (EC); (c) activates signalling pathways leading to the introduction

CC of apoptosis in EC; and/or (d) inhibits the signalling pathway required

CC for maintenance of EC viability. (I) has cytostatic, antitumour,

CC antiatherosclerotic, vasotropic, vulnary, tranquilliser, thrombolytic,

CC ophthalmological, gynaecological, antiulcer, antidiabetic, antiarthritic,

CC antiangiogenic, antiapoptotic and endocrine activities. An antibody (IX)

CC specific for an epitope of (I) is useful for inhibiting tumour growth or

CC angiogenesis in a subject. (I), a D5 fusion polypeptide (II) or a dimeric

CC or trimeric fusion polypeptide (III) can be used for inhibiting EC

CC migration, proliferation, invasion, or angiogenesis, or for inducing EC

CC apoptosis. An angiogenic EC-targeting pharmaceutical composition (X)

CC comprising (I), (II), or (III), can be used for treating a subject having

CC a disease or condition associated with undesired EC migration,

CC proliferation, invasion or angiogenesis. (I), (II), or (III) can be used

CC for isolating a D5 domain binding molecule from a complex mixture and for

CC isolating or enriching cells expressing D5 domain binding sites from a

CC cell mixture. The present sequence represents the human high molecular

CC weight kininogen (HK) protein, which is given in the exemplification of

CC the present invention

XX Sequence 644 AA;

SQ Query Match 100.0%; Score 660; DB 5; Length 644;

Best Local Similarity 100.0%; Pred. No. 2.3e-65;

Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVQPPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKARVQV 60

Db 253 GKDFVQPPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKARVQV 312

QY 61 AGKKYFDVFARETTCKESNEELTESCTKLGQSLDCNAEYVVPWEKKIYPTVNCQ 120

Db 313 AGKKYFDVFARETTCKESNEELTESCTKLGQSLDCNAEYVVPWEKKIYPTVNCQ 372

QY 121 LGM 123

Db 373 LGM 375

RESULT 13

ABU99150

ID ABU99150 standard; protein; 644 AA.

XX ABU99150;

XX 01-AUG-2003 (first entry)

XX Novel human GPCR related protein NOV12h.

DE Human; G-protein coupled receptor related protein; GPCR related protein;

KW NOV; cytostatic; cardiant; antiatherosclerotic; antidiabetic;

KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;

KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;

KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;

KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.

XX Homo sapiens.

XX WO200299116-A2.

XX PD 12-DEC-2002.

XX PF 04-JUN-2002; 2002WO-US017428.

XX PR 04-JUN-2001; 2001US-0295607P.

XX PR 04-JUN-2001; 2001US-0295661P.

XX PR 06-JUN-2001; 2001US-0296404P.

XX PR 06-JUN-2001; 2001US-0296418P.

XX PR 14-JUN-2001; 2001US-0298285P.

XX PR 15-JUN-2001; 2001US-0298556P.

XX PR 21-JUN-2001; 2001US-0299949P.

XX PR 26-JUN-2001; 2001US-0300883P.

XX PR 28-JUN-2001; 2001US-0301550P.

XX PR 13-AUG-2001; 2001US-0311972P.

XX PR 27-AUG-2001; 2001US-0315071P.

XX PR 29-AUG-2001; 2001US-0315660P.

XX PR 14-SEP-2001; 2001US-032293P.

XX PR 17-SEP-2001; 2001US-0322706P.

XX PR 14-DEC-2001; 2001US-0341186P.

XX PR 28-FEB-2002; 2002US-0361189P.

XX PR 12-MAR-2002; 2002US-0363673P.

XX PR 03-JUN-2002; 2002US-00363676.

XX (CURA-) CURAGEN CORP.

XX Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR, Li L; Gangoli EA, Gerlach VL, Gorman L, Guo X, Hjalit T, Kekuda R, Li L; MacDougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M; Pena CBA, Rastelli L, Shimkets RA, Stone DJ, Spytek KA, Vernet CAM; Voss EZ, Zerhusen BD;
 DR WPI; 2003-140627/13.
 DR N-PSDB; ACD03654.

XX New NOVX polypeptides and nucleic acids, useful for preventing or treating NOVX-associated disorders, e.g. cancer, cardiomyopathy, atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or pharmacogenomics.

XX Claim 1; Page 148; 332pp; English.

XX The invention describes an isolated polypeptide (I) comprising any of 27 118-961 residue amino acid sequences, given in the specification, a mature form of them, a sequence that is at least 95 % identical to them, or a sequence having one or more conservative substitutions in them. The polypeptide is useful in manufacturing a medicament for treating a syndrome associated with a human disease selected from a pathology associated with the polypeptide. The NOVX polypeptides, polynucleotides and antibodies are useful in treating or preventing NOVX-associated disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune disease, Alzheimer's disease, asthma, haematopoietic disorders, Parkinson's disease, AIDS, obesity, asthma, haematopoietic disorders, cancer-associated cachexia, and other wasting disorders associated with chronic diseases. The nucleic acids and polypeptides may also be used as targets for the identification of small molecules that modulate or inhibit e.g. neurogenesis, cell differentiation, cell proliferation, haematopoiesis, wound healing and angiogenesis, in gene therapy, in generation of antibodies that bind immunospecifically to NOVX substances for use in therapeutic or diagnostic methods. The nucleic acids are further used as hybridisation probes, in chromosome mapping, tissue typing, preventive medicine, and pharmacogenomics. The polypeptides are also useful as

CC vaccines. This is the amino acid sequence of a novel human G-protein coupled receptor related protein NOV

XX SQ Sequence 644 AA;

Query Match 100.0%; Score 660; DB 6; Length 644;
 Best Local Similarity 100.0%; Pred. No. 2.3e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVOPPTKICVGCPRDPTNSPELEETLTHITKLNAENNAATFYFKIDNVKKARQVYV 60
 |||||
 DB 253 GKDFVOPPTKICVGCPRDPTNSPELEETLTHITKLNAENNAATFYFKIDNVKKARQVYV 312
 |||||

QY 61 AGKYYFIDFVARETTCSKESNEELTESCTKKLGQSLDCNAEYVYVWPKKIYPTVNCQP 120
 |||||
 DB 313 AGKYYFIDFVARETTCSKESNEELTESCTKKLGQSLDCNAEYVYVWPKKIYPTVNCQP 372
 |||||

QY 121 LGM 123
 |||

DB 373 LGM 375

RESULT 14

ABU99145
 ID ABU99145 standard; protein; 644 AA.

XX AC ABU99145;

XX DT 01-AUG-2003 (first entry)

XX DE Novel human GPCR related protein NOV12c.

XX Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytostatic; cardiant; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.

XX OS Homo sapiens.

XX PN WO200299116-A2.

XX PD 12-DEC-2002.

XX PF 04-JUN-2002; 2002WO-US017428.

XX PR 04-JUN-2001; 2001US-0295607P.

XX PR 04-JUN-2001; 2001US-0295661P.

XX PR 06-JUN-2001; 2001US-0296404P.

XX PR 06-JUN-2001; 2001US-0296418P.

XX PR 14-JUN-2001; 2001US-0298285P.

XX PR 15-JUN-2001; 2001US-0298556P.

XX PR 21-JUN-2001; 2001US-0299949P.

XX PR 28-JUN-2001; 2001US-0300883P.

XX PR 13-AUG-2001; 2001US-0311972P.

XX PR 27-AUG-2001; 2001US-0315071P.

XX PR 29-AUG-2001; 2001US-0315660P.

XX PR 14-SEP-2001; 2001US-032293P.

XX PR 17-SEP-2001; 2001US-0322706P.

XX PR 14-DEC-2001; 2001US-0341186P.

XX PR 28-FEB-2002; 2002US-0361189P.

XX PR 12-MAR-2002; 2002US-0363673P.

XX PR 12-MAR-2002; 2002US-0363676P.

XX PR 03-JUN-2002; 2002US-00363676.

XX (CURA-) CURAGEN CORP.

XX Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR,
 PI Gangoli EA, Gerlach VL, Gorman L, Guo X, Hjalt T, Kekuda R, Li L,
 PI Macdougall JR, Malyankar UM, Millet I, Padigar M, Patturajan M,
 PI Pena CEA, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet CM,
 PI Voss EZ, Zerhusen BD;
 XX WPI; 2003-140627/13.
 DR N-PSDB; ACD03649.
 XX
 XX New NOVX polypeptides and nucleic acids, useful for preventing or
 PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX
 XX Claim 1; Page 144-145; 332pp; English.
 XX
 XX The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX
 XX Sequence 644 AA;

Query Match 100.0%; Score 660; DB 6; Length 644;
 Best Local Similarity 100.0%; Pred. No. 2.3e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GKDFVQPTKICVGCPRDPTNSPELEETLTHITIKLAENNAATFEKIDNVKARVQV 60
 |||||
 Db 253 GKDFVQPTKICVGCPRDPTNSPELEETLTHITIKLAENNAATFEKIDNVKARVQV 312
 |||||

Qy 61 AGKKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTVNCQP 120
 |||||
 Db 313 AGKKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTVNCQP 372
 |||||

Qy 121 LGM 123
 |||||
 Db 373 LGM 375
 |||||

RESULT 15
 AAB37447
 ID AAB37447 standard; protein; 122 AA.
 XX
 AC AAB37447;
 XX
 DT 21-FEB-2001 (first entry)
 XX
 XX Human kininogen D3.
 DE
 XX Enzyme; legumain; endopeptidase; cystatin; human; kininogen.
 KW
 XX Homo sapiens.
 OS
 XX

FN WC200064945-A1.
 XX
 PD 02-NOV-2000.
 XX
 PF 20-APR-2000; 2000WO-GB001571.
 XX
 XX 22-APR-1999; 99GB-00009133.
 PR
 XX (BABR-) BABRAHAM INST.
 XX
 XX Abrahamson M, Barrett AJ;
 XX WPI; 2000-687316/67.
 XX
 XX Inhibition of mammalian legumain or legumain-related endopeptidase by
 PT cystatin involves interaction with second papain-non-reactive site of
 PT cystatin.
 XX
 PS Disclosure; Fig 4; 45pp; English.
 XX
 XX The present invention relates to inhibition of the enzymatic activity of
 CC legumain or a legumain-related endopeptidase by cystatin. The inhibition
 CC involves an interaction between legumain and a papain-non-reactive site
 CC of cystatin. Legumain (EC 3.4.22.34) is a cysteine endopeptidase, and
 CC performs a protein-processing function. The present sequence is human
 CC kininogen D3, which was used in the present invention. Kininogen is a
 CC type 3 cystatin
 XX
 XX Sequence 122 AA;
 SQ
 Query Match 95.2%; Score 628; DB 3; Length 122;
 Best Local Similarity 100.0%; Pred. No. 9.5e-63;
 Matches 117; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 7 PPTKICVGCPRDPTNSPELEETLTHITIKLAENNAATFEKIDNVKARVQVAGKKYF 66
 |||||
 Db 1 PPTKICVGCPRDPTNSPELEETLTHITIKLAENNAATFEKIDNVKARVQVAGKKYF 60
 |||||

Qy 67 IDFVARETTCKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTVNCQPLGM 123
 |||||
 Db 61 IDFVARETTCKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTVNCQPLGM 117
 |||||

Search completed: September 24, 2004, 14:08:37
 Job time : 52.692 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:04:32 ; Search time 8.364 Seconds
(without alignments)
765.738 Million cell updates/sec

Title: US-10-661-784-1

Perfect score: 660

Sequence: 1 GKDFVQPTKICVGRDIP.....YVFWKIKYPTVNCQPLGM 123

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_42.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	660	100.0	644	1 KNG_HUMAN	P01042 homo sapien
2	477	72.3	436	1 KNL1_BOVIN	P01046 bos taurus
3	477	72.3	621	1 KKH1_BOVIN	P01044 bos taurus
4	450	68.2	434	1 KNL2_BOVIN	P01047 bos taurus
5	450	68.2	619	1 KKH2_BOVIN	P01045 bos taurus
6	430	65.2	661	1 KNG_MOUSE	O08677 mus musculus
7	426	64.5	639	1 KNG_RAT	P08934 rattus norv
8	409	62.0	430	1 KNT2_RAT	P08932 rattus norv
9	401	60.8	430	1 KNT1_RAT	P01048 rattus norv
10	175	26.5	144	1 CYTF_MOUSE	O09098 mus musculus
11	165.5	25.1	145	1 CYTF_HUMAN	O76096 mus sapien
12	138.5	21.0	149	1 CYTM_HUMAN	Q15828 homo sapien
13	136	20.6	148	1 CYTC_BOVIN	P01035 bos taurus
14	133.5	20.2	146	1 CYTC_MACMU	O19092 macaca mula
15	132.5	20.1	146	1 CYTC_SALIC	O19093 salmieri sci
16	130	19.7	127	1 CYTC_RAT	P14841 rattus norv
17	129.5	19.6	140	1 CYTC_MOUSE	P21460 mus musculus
18	129.5	19.6	378	1 FETB_RAT	Q9QX79 rattus norv
19	129.5	19.6	382	1 FETB_HUMAN	Q9UGM5 homo sapien
20	128	19.4	111	1 CYT_BITAR	P08935 bitis ariet
21	127.5	19.3	146	1 CYTC_HUMAN	P01034 homo sapien
22	125	18.9	141	1 CYTT_HUMAN	P09228 homo sapien
23	124.5	18.9	148	1 CYTC_RABIT	O07862 oryctolagus
24	122.5	18.6	116	1 CYT_COTJA	P10661 coturnix co
25	118.5	18.0	139	1 CYT_CHICK	P01038 gallus gall
26	115.5	17.5	388	1 FETB_MOUSE	Q9QXCI mus musculus
27	113.5	17.2	122	1 CYTA_SARPE	P31727 sarcophaga
28	113	17.1	141	1 CYTS_RAT	P19313 rattus norv
29	109.5	16.6	141	1 CYTN_HUMAN	P01037 homo sapien
30	109	16.5	141	1 CYTS_HUMAN	P01036 homo sapien
31	108.5	16.4	130	1 CYT_ONCKE	Q38967 oncorhynch
32	108	16.4	129	1 CYT_CYPCA	P35481 cyprius ca
33	106.5	16.1	130	1 CYT_ONCMY	Q91195 oncorhynch

ALIGNMENTS

RESULT 1

KNG_HUMAN
ID KNG_HUMAN STANDARD; PRT; 644 AA.
AC P01042; P01043;
DT 21-JUL-1986 (Rel. 01, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Kininogen precursor (Alpha-2-thiol proteinase inhibitor) [Contains: Bradykinin]
DE KNG.
GN KNG.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OC NCBI_TaxID=9606;
RX KNG [1]
RP SEQUENCE FROM N.A. (ISOFORMS HMW AND LMW).
RC TISSUE=Liver;
RX MEDLINE=85234582; PubMed=2989293;
RA Takagaki Y., Kitamura N., Nakanishi S.;
RT "Cloning and sequence analysis of cDNAs for human high molecular weight and low molecular weight prekininogens. Primary structures of two human prekininogens.";
RT J. Biol. Chem. 260:8601-8609(1985).
RN [2]
RP GENE STRUCTURE.
RX MEDLINE=85234583; PubMed=2989294;
RA Kitamura N., Kitagawa H., Fukushima D., Takagaki Y., Miyata T., Nakanishi S.;
RT "Structural organization of the human kininogen gene and a model for its evolution.";
RT J. Biol. Chem. 260:8610-8617(1985).
RN [3]
RP SEQUENCE OF 1-401 FROM N.A.
RX MEDLINE=85122621; PubMed=6441591;
RA Ohkubo I., Kurachi K., Takasawa T., Shiohara H., Sasaki M.;
RT "Isolation of a human cDNA for alpha 2-thiol proteinase inhibitor and its identity with low molecular weight kininogen.";
RL Biochemistry 23:5691-5697(1984).
RN [4]
RP SEQUENCE OF 379-644.
RX MEDLINE=86030270; PubMed=4054110;
RA Lottspeich F., Kellermann J., Henschen A., Foersts B., Mueller-Esterl W.;
RT "The amino acid sequence of the light chain of human high-molecular-mass kininogen.";
RL Eur. J. Biochem. 152:307-314(1985).
RN [5]
RP SEQUENCE OF 381-389.
RX MEDLINE=90255622; PubMed=4952632;
RA Pierce J.V.;
RT "Structural features of plasma kinins and kininogens.";
RL Fed. Proc. 27:52-57(1968).
RN [6]
RP DISULFIDE BONDS.
RA Sueyoshi T., Miyata T., Kato H., Iwanaga S.;
RT "Disulfide bonds in bovine HMW kininogens.";

P22085 onchocerca
Q9d269 mus musculus
Q9h114 homo sapien
P32766 mus musculus
Q9h112 homo sapien
P29699 mus musculus
P81714 naja atra (m
P31726 zea mays (m
P28325 homo sapien
P24090 rattus norv
O60676 homo sapien
O88969 rattus norv

34 105.5 16.0 162 1 CYTX_ONCVO
35 101 15.3 139 1 CS11_MOUSE
36 95 14.4 165 1 CS11_HUMAN
37 94.5 14.3 142 1 CST8_MOUSE
38 93.5 14.2 137 1 CS11_HUMAN
39 93.5 14.2 345 1 A2HS_MOUSE
40 93 14.1 99 1 CYT_NAJAT
41 91.5 13.9 135 1 CYT1_MAIZE
42 91 13.8 142 1 CYTD_HUMAN
43 90.5 13.7 352 1 A2HS_RAT
44 89.5 13.6 142 1 CST8_HUMAN
45 86.5 13.1 142 1 CST8_RAT

Seikagaku 56:808-808(1984).
 [7]
 RY CARBOHYDRATE-LINKAGE SITE ASN-294.
 RX MEDLINE=22660472; PubMed=12754519;
 RA Zhang H., Li X.-J., Martin D.B., Abersold R.;
 RT "Identification and quantification of N-linked glycoproteins using
 RT hydrazide chemistry, stable isotope labeling and mass spectrometry.";
 RL Nat. Biotechnol. 21:660-666(2003).
 CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 CC HMW-kininogen plays an important role in blood coagulation by
 CC helping to position optimally prekallikrein and factor XI next to
 CC factor XII; (3) HMW-kininogen inhibits the thrombin-and plasmin-
 CC induced aggregation of thrombocytes; (4) the active peptide
 CC bradykinin that is released from HMW-kininogen shows a variety of
 CC physiological effects: (4A) influence in smooth muscle
 CC contraction, (4B) induction of hypotension, (4C) natriuresis and
 CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
 CC mediator of inflammation and causes (4E1) increase in vascular
 CC permeability, (4E2) stimulation of nociceptors (4E3) release of
 CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
 CC a cardioprotective effect (directly via bradykinin action,
 CC indirectly via endothelium-derived relaxing factor action); (5)
 CC LMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
 CC kininogen is in contrast to HMW-kininogen not involved in blood
 CC clotting.
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- ALTERNATIVE PRODUCTS.
 CC Event=Alternative splicing; Named isoforms=2;
 CC Name=HMW;
 CC IsoId=P01042-1; Sequence=Displayed;
 CC Name=LMW;
 CC IsoId=P01042-2; Sequence=VSP_001261, VSP_001262;
 CC -!- TISSUE SPECIFICITY: Plasma.
 CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 CC -!- SIMILARITY: Contains 3 cystatin-like domains.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; K02566; AAA35497.1; -;
 DR EMBL; M11437; AAB59550.1; JOINED.
 DR EMBL; M11438; AAB59550.1; JOINED.
 DR EMBL; M11521; AAB59550.1; JOINED.
 DR EMBL; M11522; AAB59550.1; JOINED.
 DR EMBL; M11523; AAB59550.1; JOINED.
 DR EMBL; M11524; AAB59550.1; JOINED.
 DR EMBL; M11525; AAB59550.1; JOINED.
 DR EMBL; M11526; AAB59550.1; JOINED.
 DR EMBL; M11527; AAB59550.1; JOINED.
 DR EMBL; M11528; AAB59550.1; JOINED.
 DR EMBL; M11529; AAB59550.1; JOINED.
 DR EMBL; M11530; AAB59550.1; JOINED.
 DR EMBL; M11437; AAB59551.1; -;
 DR EMBL; M11438; AAB59551.1; JOINED.
 DR EMBL; M11521; AAB59551.1; JOINED.
 DR EMBL; M11522; AAB59551.1; JOINED.
 DR EMBL; M11523; AAB59551.1; JOINED.
 DR EMBL; M11524; AAB59551.1; JOINED.
 DR EMBL; M11525; AAB59551.1; JOINED.
 DR EMBL; M11526; AAB59551.1; JOINED.
 DR EMBL; M11527; AAB59551.1; JOINED.
 DR EMBL; M11528; AAB59551.1; JOINED.
 DR PIR; A01279; KGHU1.
 DR PIR; A01280; KGHU1.
 DR SWISS-2DPAGE; P01042; HUMAN.
 DR Genew; HGNC:6383; KNG.
 DR MIM; 228960; -;
 DR GO; GO:0007596; P:blood coagulation; NAS.
 DR GO; GO:0030146; P:diuresis; NAS.
 DR GO; GO:0006954; P:inflammatory response; NAS.

DR GO; GO:0030147; P:natriuresis; NAS.
 DR GO; GO:0006939; P:smooth muscle contraction; NAS.
 DR InterPro; IPR000010; Cystatin.
 DR InterPro; IPR002395; Kininogen.
 DR Pfam; PFO0031; cystatin; 3.
 DR PRINTS; PR000334; KININOGEN.
 DR SMART; SM00043; CY; 3.
 DR PROSITE; PS00287; CYSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Thiol protease inhibitor; Vasodilator;
 KW Bradykinin; Blood coagulation; Inflammatory response; Signal;
 KW Alternative splicing; Pyrrolidone carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 644 KININOGEN.
 FT CHAIN 19 380 KININOGEN HEAVY CHAIN.
 FT PEPTIDE 381 389 BRADYKININ.
 FT CHAIN 390 644 KININOGEN LIGHT CHAIN.
 FT DOMAIN 19 136 CYSTATIN-LIKE 1.
 FT DOMAIN 137 258 CYSTATIN-LIKE 2.
 FT DOMAIN 259 380 CYSTATIN-LIKE 3.
 FT DOMAIN 420 510 HIS-RICH
 FT (ASSOCIATED WITH CLOTTING ACTIVITY).
 FT REPEAT 420 449
 FT REPEAT 450 479
 FT REPEAT 480 510
 FT MOD RES 19 19
 FT DISULFID 28 614
 FT DISULFID 83 94
 FT DISULFID 107 126
 FT DISULFID 142 145
 FT DISULFID 206 218
 FT DISULFID 229 248
 FT DISULFID 264 267
 FT DISULFID 328 340
 FT DISULFID 351 370
 FT CARBOHYD 48 48
 FT CARBOHYD 169 169
 FT CARBOHYD 205 205
 FT CARBOHYD 294 294
 FT CARBOHYD 401 401
 FT CARBOHYD 533 533
 FT CARBOHYD 542 542
 FT CARBOHYD 546 546
 FT CARBOHYD 557 557
 FT CARBOHYD 571 571
 FT CARBOHYD 577 577
 FT CARBOHYD 593 593
 FT CARBOHYD 628 628
 FT VARSPLIC 402 427
 FT VARSPLIC 428 644
 FT CONFLICT 593 593
 FT SEQUENCE 644 AA; 71945 MW; 313284CBFA8FBB7E CRC64;
 Query Match 100.0%; Score 660; DB 1; Length 644;
 Best Local Similarity 100.0%; Pred. No. 2e-55; 0; Gaps 0;
 Matches 123; Conservative 0; Mismatches 0; Indels 0;
 Qy 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHTTKLNAENNAATFYFKIDNVKARQVQV 60
 Db 253 GKDFVQPTKICVGCPRDIPNTSPLEETLTHTTKLNAENNAATFYFKIDNVKARQVQV 312
 Qy 61 AGKYSIDFVARETTCKSKESNELTSCCTKKLGQSLDCNAEYVVVWPKKIYPTVNCOP 120
 Db 313 AGKYSIDFVARETTCKSKESNELTSCCTKKLGQSLDCNAEYVVVWPKKIYPTVNCOP 372
 Qy 121 LGM 123
 Db 373 LGM 375
 RESULT 2

KNL1_BOVIN
 ID KNL1_BOVIN STANDARD; PRT; 436 AA.
 AC P01046;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Kininogen, LMW I precursor (Thiol proteinase inhibitor) [Contains:
 Bradykinin]
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovoidea;
 OC Bovidae; Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=83117859; PubMed=6572010;
 RA Nawa H., Kitamura N., Hirose T., Asai M., Inayama S., Nakanishi S.;
 RT "Primary structures of bovine liver low molecular weight kininogen
 precursors and their two mRNAs.";
 RL Proc. Natl. Acad. Sci. U.S.A. 80:90-94(1983).
 RN [2]
 RP SEQUENCE OF 19-378.
 RX MEDLINE=87137530; PubMed=3546295;
 RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
 RA Miyata T., Iwanaga S.;
 RT "Bovine high molecular weight kininogen. The amino acid sequence,
 positions of carbohydrate chains and disulfide bridges in the heavy
 chain portion.";
 RL J. Biol. Chem. 262:2768-2779(1987).
 CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 LMW-kininogen inhibits the aggregation of thrombocytes; (3) the
 active peptide kallidin that is released from LMW-kininogen shows
 a variety of physiological effects: (3A) influence in smooth
 muscle contraction, (3B) induction of hypotension, (3C)
 natriuresis and diuresis (kidney).
 CC -!- SUBCELLULAR LOCATION: Extracellular.
 CC -!- ALTERNATIVE PRODUCTS:
 Event=Alternative splicing; Named isoforms=2;
 Name=LMW I;
 Name=HMW I;
 IsoId=P01046-1; Sequence=Displayed;
 IsoId=P01044-1; Sequence=External;
 CC -!- TISSUE SPECIFICITY: Plasma.
 CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 CC -!- MISCELLANEOUS: LMW-kininogen is in contrast to HMW-kininogen not
 involved in blood clotting.
 CC -!- SIMILARITY: Contains 3 cystatin-like domains.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 between the Swiss Institute of Bioinformatics and the EMBL outstation -
 the European Bioinformatics Institute. There are no restrictions on its
 use by non-profit institutions as long as its content is in no way
 modified and this statement is not removed. Usage by and for commercial
 entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 or send an email to license@sb-sib.ch).
 CC
 DR EMBL; V00426; CAA23709.1; -;
 DR PIR; A01283; KGBOL1.
 DR InterPro; IP3000010; Cystatin.
 DR Pfam; PF00031; cystatin; 3.
 DR SMART; SM00043; Cy; 3.
 DR PROSITE; PS00287; CYSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;
 KW Thiol protease inhibitor; Bradykinin; Signal;
 KW Pyrrolidone carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 436 KININOGEN, LMW I.
 FT CHAIN 19 378 HEAVY CHAIN.
 FT PEPTIDE 380 388 BRADYKININ.
 FT CHAIN 389 436 LIGHT CHAIN.
 FT DOMAIN 19 135 CYSTATIN-LIKE 1.
 FT DOMAIN 136 257 CYSTATIN-LIKE 2.
 FT DOMAIN 258 378 CYSTATIN-LIKE 3.

FT MOD RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
 FT CARBOHYD 87 87 N-LINKED (GLCNAC. . .).
 FT CARBOHYD 136 136 O-LINKED (PARTIAL).
 FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
 FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
 FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
 FT DISULFID 27 406 INTERCHAIN.
 FT DISULFID 82 93
 FT DISULFID 106 125
 FT DISULFID 141 144
 FT DISULFID 205 217
 FT DISULFID 228 247
 FT DISULFID 263 286
 FT DISULFID 327 339
 FT DISULFID 350 369
 FT CONFLICT 295 295 A -> T (IN REF. 1; CAA23709).
 SQ SEQUENCE 436 AA; 48427 MW; F01F7EB6814BCE6C CRC64;
 Query Match 72.3%; Score 477; DB 1; Length 436;
 Best Local Similarity 71.9%; Pred. No. 3.6e-38;
 Matches 87; Conservative 14; Mismatches 20; Indels 0; Gaps 0;
 QY 2 KDFVQPTKICVGCPRDIPTNSPELEETLTHITKLNNAENNAFFPKIDNVKKARQVVA 61
 DB 253 KDFVQPTKLCAGCPKIPVDSFDLEELSHSIAKLNAEHDGAFYFKIDTVKKATQVVA 312
 QY 62 GKRYFIDFVARETTCSENEELTSCETPKLGQSLDCNAEYVVPWEKKIYPTVNCQPL 121
 DB 313 GLKYSIVFIARETTCSENEELTSCSEINIRHQILHCDANVYVVPWEKKYPTVNCQPL 372
 QY 122 G 122
 DB 373 G 373
 RESULT 3
 KNL1_BOVIN STANDARD; PRT; 621 AA.
 ID KNL1_BOVIN
 AC P01044;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Kininogen, HMW I precursor (Thiol proteinase inhibitor) [Contains:
 Bradykinin].
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovoidea;
 OC Bovidae; Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=84014106; PubMed=6571699;
 RA Kitamura N., Takagaki Y., Furuto S., Tanaka T., Nawa H., Nakanishi S.;
 RT "A single gene for bovine high molecular weight and low molecular
 weight kininogens.";
 RL Nature 305:545-549(1983).
 RN [2]
 RP SEQUENCE OF 19-378.
 RX MEDLINE=87137530; PubMed=3546295;
 RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
 RA Miyata T., Iwanaga S.;
 RT "Bovine high molecular weight kininogen. The amino acid sequence,
 positions of carbohydrate chains and disulfide bridges in the heavy
 chain portion.";
 RL J. Biol. Chem. 262:2768-2779(1987).
 RN [3]
 RP SEQUENCE OF 378-393.
 RX MEDLINE=70180420; PubMed=4986212;
 RA Kato H., Nagasawa S., Suzuki T.;
 RT "Studies on the structure of bovine kininogen: cleavages of disulfide
 bonds and of methionyl bonds in kininogen-II.";
 RL J. Biochem. 67:313-323(1970).
 RN [4]

RP SEQUENCE OF 458-498.
RX MEDLINE=75170265; PubMed=1169237;
RA Han Y N., Komiya M., Iwanaga S., Suzuki T.;
RT "Studies on the primary structure of bovine high-molecular-weight
RT kininogen. Amino acid sequence of a fragment ('histidine-rich
RT peptide') released by plasma kallikrein.";
RL J. Biochem. 77:55-68(1975).
CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC HMW-kininogen plays an important role in blood coagulation by
CC helping to position optimally prekallikrein and factor XI next to
CC factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
CC induced aggregation of thrombocytes; (4) the active peptide
CC bradykinin that is released from HMW-kininogen shows a variety of
CC physiological effects: (4A) influence in smooth muscle
CC contraction, (4B) induction of hypotension, (4C) natriuresis and
CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
CC mediator of inflammation and causes (4E1) increase in vascular
CC permeability, (4E2) stimulation of nociceptors (4E3) release of
CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
CC a cardioprotective effect (directly via bradykinin action,
CC indirectly via endothelium-derived relaxing factor action).
CC -!- SUBCELLULAR LOCATION: Extracellular.
CC -!- ALTERNATIVE PRODUCTS:
CC Event-Alternative splicing; Named isoforms=2;
CC Name=HMW I;
CC IsoId=P01044-1; Sequence=Displayed;
CC IsoId=P01046-1; Sequence=External;
CC -!- TISSUE SPECIFICITY: Plasma.
CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -!- SIMILARITY: Contains 3 cystatin-like domains.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC
CC -----
CC EMBL: V01491; CAA24735.1; -.
CC PIR: A01281; KGBOH1.
CC InterPro: IPR000010; Cystatin.
CC InterPro: IPR002395; Kininogen.
CC Pfam: PF00031; Cystatin_3.
CC PRINTS: PRO0334; KININOGEN.
CC SMART: SM00043; CY; 3.
CC PROSITE: PS00287; CYSTATIN; 2.
CC Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;
CC Thiol protease inhibitor; Bradykinin; Blood coagulation;
CC Inflammatory response; Signal; Pyrrolidone carboxylic acid.
CC SIGNAL 1 18 PROBABLE
CC CHAIN 19 621 KININOGEN, HMW I.
CC CHAIN 19 378 HEAVY CHAIN.
CC PEPTIDE 380 388 BRADYKININ.
CC CHAIN 389 621 LIGHT CHAIN.
CC DOMAIN 19 135 CYSTATIN-LIKE 1.
CC DOMAIN 136 257 CYSTATIN-LIKE 2.
CC DOMAIN 258 378 CYSTATIN-LIKE 3.
CC MOD RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
CC CARBOHYD 87 87 N-LINKED (GLCNAC. . .).
CC CARBOHYD 136 136 O-LINKED (PARTIAL).
CC CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
CC CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
CC CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
CC DISULFID 27 591 INTERCHAIN.
CC DISULFID 82 93
CC DISULFID 106 125
CC DISULFID 141 144
CC DISULFID 205 217
CC DISULFID 228 247
CC DISULFID 263 266
CC DISULFID 327 339

FT DISULFID 350 369
SQ SEQUENCE 621 AA; 68890 MW; D16850BEF3C55CD CRC64;
Query Match 72.3%; Score 477; DB 1; Length 621;
Best Local Similarity 71.9%; Pred. No. 5.4e-38;
Matches 87; Conservative 14; Mismatches 20; Indels 0; Gaps 0;
QY 2 KDFVQPTKICVGCPRDIPDTPNSPELEETLTHITKLAENNAFFPKIDNVKARVQVVA 61
DB 253 KDFVQPTKICVGCPRDIPDTPNSPELEETLTHITKLAENNAFFPKIDNVKARVQVVA 312
QY 62 GKQYPTDFVARETTCSENEELTSCSKLQSLDCNAEVVVPWEKKIYPTVNCQPL 121
DB 313 GLKISIVFIARETTCSENEELTSCSKLQSLDCNAEVVVPWEKKIYPTVNCQPL 372
QY 122 G 122
DB 373 G 373
RESULT 4
KML2 BOVIN STANDARD; PRT; 434 AA.
ID KML2 BOVIN
AC P01047;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Kininogen, LMW II precursor (Thiol proteinase inhibitor) [Contains:
DE Bradykinin].
DE Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Euthera; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=83117859; PubMed=6572010;
RA Nawa H., Kitamura N., Hirose T., Asai M., Inayama S., Nakanishi S.;
RT "Primary structures of bovine liver low molecular weight kininogen
RT precursors and their two mRNAs";
RL Proc. Natl. Acad. Sci. U.S.A. 80:90-94 (1983).
RN [2]
RP SEQUENCE OF 19-376..
RX MEDLINE=87137530; PubMed=3546285;
RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
RA Miyata T., Iwanaga S.;
RT "Bovine high molecular weight kininogen. The amino acid sequence,
RT positions of carbohydrate chains and disulfide bridges in the heavy
RT chain portion.";
RL J. Biol. Chem. 262:2768-2779 (1987).
CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC LMW-kininogen inhibits the aggregation of thrombocytes; (3) the
CC active peptide kallidin that is released from LMW-kininogen shows
CC a variety of physiological effects: (3A) influence in smooth
CC muscle contraction, (3B) induction of hypotension, (3C)
CC natriuresis and diuresis (kidney).
CC -!- SUBCELLULAR LOCATION: Extracellular.
CC -!- ALTERNATIVE PRODUCTS:
CC Event-Alternative splicing; Named isoforms=2;
CC Name=LMW II;
CC IsoId=P01047-1; Sequence=Displayed;
CC Name=HMW II;
CC IsoId=P01045-1; Sequence=External;
CC -!- TISSUE SPECIFICITY: Plasma.
CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -!- MISCELLANEOUS: LMW-kininogen is in contrast to HMW-kininogen not
CC involved in blood clotting.
CC -!- SIMILARITY: Contains 3 cystatin-like domains.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way

CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).

```

DR EMBL; V00427; CAA23710.1; -.
DR PIR; A01284; KGBOL2.
DR HSP; P01038; IA90.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 3.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;
KW Thiol protease inhibitor; Bradykinin; Signal;
KW Pyrrolidone carboxylic acid.
FT SIGNAL 1 18
FT CHAIN 19 434 KININOGEN, LMW II.
FT PEPTIDE 19 376 HEAVY CHAIN.
FT CHAIN 19 376 BRADYKININ.
FT CHAIN 378 386 BRADYKININ.
FT CHAIN 387 434 LIGHT CHAIN.
FT DOVAIN 19 135 CYSTATIN-LIKE 1.
FT DOVAIN 136 256 CYSTATIN-LIKE 2.
FT DOVAIN 257 376 CYSTATIN-LIKE 3.
FT MOD RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
FT CARBOHYD 87 87 O-LINKED (GLCNAC. . .).
FT CARBOHYD 136 136 O-LINKED (PARTIAL).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
FT CARBOHYD 280 280 N-LINKED (GLCNAC. . .).
FT DISULFID 27 404 INTERCHAIN.
FT DISULFID 82 93
FT DISULFID 106 125
FT DISULFID 141 144
FT DISULFID 205 217
FT DISULFID 228 247
FT DISULFID 261 264
FT DISULFID 325 337
FT DISULFID 348 367
SQ SEQUENCE 434 AA; 48148 MW; 73A07079DE3E03430 CRC64;

Query Match 68.2%; Score 450; DB 1; Length 434;
Best Local Similarity 68.9%; Pred. NO. 1.4e-35;
Matches 84; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

OY 1 GKDFVQPTKICVGPDPITNSPELETHTITKLNAENNATFYKIDNVKKARQVQV 60
DB 252 GEDEL--PPMVCVCPKIPVDSPDLSEALNHSIAKLNAEHDGFYKIDTVKATQVQV 309

OY 61 AGKYFIDFVARETTCSENEELTESCETKKLQSLDCNAEVVYVPWEKKIYPTVNCQP 120
DB 310 GGLKYSIVFIARETTCSENEELTKSCENIHQILHCDANVYVPWEKKVYPTVNCQP 369

OY 121 LG 122
DB 370 LG 371

RESULT 5
KNH2_BOVIN
ID_KNH2_BOVIN STANDARD; PRT; 619 AA.
AC P01045;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Kininogen, HMW II precursor (Thiol proteinase inhibitor) [Contains:
DE Bradykinin].
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.

```

```

RX MEDLINE=84014106; PubMed=6571699;
RA Kitamura N., Takagaki Y., Furuto S., Tanaka T., Nawa H., Nakanishi S.;
RT "A single gene for bovine high molecular weight and low molecular
RT weight kininogens";
RL Nature 305:545-549(1983).
RN [2]
RP SEQUENCE OF 19-376.
RX MEDLINE=87137530; PubMed=3546295;
RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
RA Miyata T., Iwanaga S.;
RT "Bovine high molecular weight kininogen. The amino acid sequence,
RT positions of carbohydrate chains and disulfide bridges in the heavy
RT chain portion.";
RL J. Biol. Chem. 262:2768-2779(1987).
RN [3]
RP SEQUENCE OF 376-391.
RX MEDLINE=70180420; PubMed=4986212;
RA Kato H., Nagasawa S., Suzuki T.;
RT "Studies on the structure of bovine kininogen: cleavages of disulfide
RT bonds and of methionyl bonds in kininogen-II.";
RL J. Biochem. 67:313-323(1970).
RN [4]
RP SEQUENCE OF 387-455.
RX MEDLINE=76260155; PubMed=956151;
RA Han Y.N., Kato H., Iwanaga S., Suzuki T.;
RT "Primary structure of bovine plasma high-molecular-weight kininogen.
RT The amino acid sequence of a glycopeptide portion (fragment 1)
RT following the C-terminus of the bradykinin moiety.";
RL J. Biochem. 79:1201-1222(1976).
RN [5]
RP SEQUENCE OF 456-496.
RX MEDLINE=75170265; PubMed=1169237;
RA Han Y.N., Komiya M., Iwanaga S., Suzuki T.;
RT "Studies on the primary structure of bovine high-molecular-weight
RT kininogen. Amino acid sequence of a fragment ('histidine-rich
RT peptide') released by plasma kallikrein.";
RL J. Biochem. 77:55-68(1975).
CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC HMW-kininogen plays an important role in blood coagulation by
CC helping to position optimally prekallikrein and factor XI next to
CC factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
CC induced aggregation of thrombocytes; (4) the active peptide
CC bradykinin that is released from HMW-kininogen shows a variety of
CC physiological effects: (4A) influence in smooth muscle
CC contraction, (4B) induction of hypotension, (4C) natriuresis and
CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
CC mediator of inflammation and causes (4E1) increase in vascular
CC permeability, (4E2) stimulation of nociceptors (4E3) release of
CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
CC a cardioprotective effect (directly via bradykinin action,
CC indirectly via endothelium-derived relaxing factor action).
CC -!- SUBCELLULAR LOCATION: Extracellular.
CC -!- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=HMW II;
CC IsoId=P01045-1; Sequence=Displayed;
CC Name=LMW II;
CC IsoId=P01047-1; Sequence=External;
CC -!- TISSUE SPECIFICITY: Plasma.
CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -!- SIMILARITY: Contains 3 cystatin-like domains.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).

```

DR InterPro: IPR000010; Cystatin.
 DR InterPro: IPR002395; Kininogen.
 DR Pfam: PF00031; Cystatin; 3.
 DR PRINTS: PRO0334; KININOGEN.
 DR SMART: SM00043; CY; 3.
 DR PROSITE: PS00287; CYSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;
 KW Thiol protease inhibitor; Bradykinin; Blood coagulation; Signal;
 KW Inflammatory response; Pyrrolidone carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 619 KININOGEN, HWV II.
 FT CHAIN 19 376 HEAVY CHAIN.
 FT PEPTIDE 378 386 BRADYKININ.
 FT CHAIN 387 619 LIGHT CHAIN.
 FT DOMAIN 19 135 CYSTATIN-LIKE 1.
 FT DOMAIN 136 256 CYSTATIN-LIKE 2.
 FT DOMAIN 257 376 CYSTATIN-LIKE 3.
 FT MOD_RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
 FT CARBOHYD 87 97 N-LINKED (GLCNAC. . .).
 FT CARBOHYD 136 136 O-LINKED (PARTIAL. . .).
 FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
 FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
 FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
 FT CARBOHYD 280 280 N-LINKED (GLCNAC. . .).
 FT CARBOHYD 400 400 O-LINKED.
 FT DISULFID 27 589 INTERCHAIN.
 FT DISULFID 82 93
 FT DISULFID 106 125
 FT DISULFID 141 144
 FT DISULFID 205 217
 FT DISULFID 228 247
 FT DISULFID 261 284
 FT DISULFID 325 337
 FT DISULFID 348 367
 FT VARIANT 398 398
 FT VARIANT 401 401
 FT VARIANT 454 454
 SQ SEQUENCE 619 AA; 68710 MW; F04320A8EB0E50DA CRC64;
 Query Match 68.2%; Score 450; DB 1; Length 619;
 Best Local Similarity 68.9%; Pred. No. 28-35;
 Matches 84; Conservative 14; Mismatches 22; Indels 2; Gaps 1;
 QY 1 GQDFVPPKICVGPDRDPTNSPELEETLTITIKLAENNATFYKIDNVKKARVQV 60
 Db 252 GDFL--PMWCVGCPKIPVDSPLDEALNHSIAKLAHDCGTFFKIDTVKKATVQV 309
 QY 61 AGKKYFIDFVARETTCKSESNEELTESCTKGLGSLDCAAFVYVVPWEKKIYPTVNCQP 120
 Db 310 GGLKYSIVFIARETTCKSGSNEELTKSCBINTHGQILHCDAVYVVPWEKKYPTVNCQP 369
 QY 121 LG 122
 Db 370 LG 371
 RESULT 6
 ID KNG_MOUSE STANDARD; PRT; 661 AA.
 AC O08677; O08676; Q91XK5;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Kininogen precursor [Contains: Bradykinin].
 GN KNG.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID:10090;
 RN [1]
 RP SEQUENCE FROM N.A. (ISOFORMS HWV AND LMW).
 RC STRAIN=C57BL/6 X CBA; TISSUE=Liver;
 RX MEDLINE=97342556; PubMed=9199253;
 Takano M., Kondo J., Yayama K., Otani M., Sano K., Okamoto H.;
 "Molecular cloning of cDNAs for mouse low-molecular-weight and high-
 molecular-weight prekininogens";
 Biochim. Biophys. Acta 1352:222-230(1997).
 [2]
 RN SEQUENCE FROM N.A. (ISOFORM LMW).
 RP STRAIN=C57BL/6J; TISSUE=Placenta;
 RC MEDLINE=22354683; PubMed=1246851;
 RX Okazaki Y., Furuno M., Kasukawa T., Adachi H., Yamanaka I., Kiyosawa H.,
 Nikaido I., Osato N., Saito R., Suzuki H., Schonbach C., Gojobori T.,
 Yagi K., Tomaru Y., Hasegawa Y., Nogami A., Quackenbush J.,
 Baldarelli R., Hill D.P., Bult C., Hume D.A., Quackenbush J.,
 Schriml L.M., Karspin A., Matsuda H., Batalov S., Beisel K.W.,
 Blake J.A., Bradt D., Brusic V., Chothia C., Corbani L.E., Cousins S.,
 Dalla E., Dragani T.A., Fletcher C.F., Forrest A., Frazer K.S.,
 Gaasterland T., Gariboldi M., Gissi C., Godzik A., Gough J.,
 Grimmond S., Gustincich S., Hirokawa N., Jackson I.J., Jarvis E.D.,
 Kanai A., Kawaji H., Kawasawa Y., Kedzierski R.M., King B.L.,
 Konagaya A., Kurochkin I.V., Lee Y., Lenhard B., Lyons P.A.,
 Maglott D.R., Maltais L., Marchionni L., McKenzie L., Miki H.,
 Nagashima T., Numata K., Okido T., Pavan W.J., Pertea G., Pesole G.,
 Petrovsky N., Pillai R., Pontius J.U., Qi D., Ramachandran S.,
 Ravasi T., Reed J.C., Reed D.J., Reid J., Ring B.Z., Ringwald M.,
 Sandelin A., Schneider C., Semple C.A., Setou M., Shimada K.,
 Sultana R., Takenaka Y., Taylor M.S., Teasdale R.D., Tomita M.,
 Verardo R., Wagner L., Wahlestedt C., Wang Y., Watanabe Y., Wells C.,
 Wilming L.G., Wynshaw-Boris A., Yanagisawa M., Yang I., Yang L.,
 Yuan Z., Zavolan M., Zhu Y., Zimmer A., Carninci P., Hayatsu N.,
 Hirozane-Kishikawa T., Kono H., Nakamura M., Sakazume N., Sato K.,
 Shiraki T., Waki K., Kawai J., Aizawa K., Arakawa T., Fukuda S.,
 Hara A., Hashizume W., Imotani K., Ishii Y., Itoh M., Kagawa I.,
 Miyazaki A., Sakai K., Sasaki D., Shibata K., Shinagawa A.,
 Yasunishi A., Yoshino M., Waterston R., Lander E.S., Rogers J.,
 Birney E., Hayashizaki Y.;
 "Analysis of the mouse transcriptome based on functional annotation of
 60,770 full-length cDNAs";
 Nature 420:563-573(2002).
 [3]
 RN SEQUENCE FROM N.A. (ISOFORM LMW).
 RP TISSUE=Liver;
 RC MEDLINE=22388257; PubMed=12477932;
 RX Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G., Schuler G.D.,
 Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 Scapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 Brownstein M.J., Ustin T.B., Toshiyuki S., Carninci P., Prange C.,
 Raha S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
 Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 Richards S., Worley K.C., Hale S.C., Garcia A.M., Gay L.J., Hulyk S.W.,
 Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 Fahey J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
 Whitting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 Butterfield Y.S.N., Krzywicki M.I., Skalska U., Smalish D.E.,
 Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 "Generation and initial analysis of more than 15,000 full-length
 human and mouse cDNA sequences";
 Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 HWV-kininogen plays an important role in blood coagulation by
 helping to position optimally prekallikrein and factor XI next to
 factor XII; (3) HWV-kininogen inhibits the thrombin and plasmin-
 induced aggregation of thrombocytes; (4) the active peptide
 bradykinin that is released from HWV-kininogen shows a variety of
 physiological effects: (4A) influence in smooth muscle
 contraction, (4B) induction of hypotension, (4C) natriuresis and
 diuresis, (4D) decrease in blood glucose level, (4E) it is a
 mediator of inflammation and causes (4E1) increase in vascular
 permeability, (4E2) stimulation of nociceptors (4E3) release of
 other mediators of inflammation (e.g. prostaglandins), (4F) it has

CC a cardioprotective effect (directly via bradykinin action,
CC indirectly via endothelium-derived relaxing factor action); (5)
CC LMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
CC kininogen is in contrast to HMW-kininogen not involved in blood
CC clotting (By similarity).
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- ALTERNATIVE PRODUCTS:
CC Event-Alternative splicing; Named isoforms=2;
CC Name=HMW;
CC IsoId=008677-1; Sequence=Displayed;
CC Name=LMW;
CC IsoId=008677-2; Sequence=VSP_001263, VSP_001264;
CC -!- TISSUE SPECIFICITY: Plasma.
CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -!- SIMILARITY: Contains 3 cysteine-like domains.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; D84435; BAA19742.1; -;
DR EMBL; D84415; BAA19742.1; -;
DR EMBL; AK005547; BAB24115.1; -;
DR EMBL; BC018158; AAB18158.1; -;
DR MGB; MGI:1097705; Kug.
DR InterPro; IPR000010; Cystatin.
DR InterPro; IPR002395; Kininogen.
DR Pfam; PF00031; cystatin; 3.
DR PRINTS; PRO0334; KININOGEN.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 1.
KW Glycoprotein; Plasma; Repeat; Thiol protease inhibitor; Vasodilator;
KW Bradykinin; Blood coagulation; Inflammatory response; Signal;
KW Alternative splicing.
FT SIGNAL 1 18
FT CHAIN 19 661 KININOGEN.
FT CHAIN 19 379 KININOGEN HEAVY CHAIN.
FT CHAIN 380 388 BRADYKININ.
FT CHAIN 389 661 KININOGEN LIGHT CHAIN.
FT DOMAIN 19 135 CYSTATIN-LIKE 1.
FT DOMAIN 136 257 CYSTATIN-LIKE 2.
FT DOMAIN 258 379 CYSTATIN-LIKE 3.
FT DOMAIN 439 524 HIS-RICH.
FT DISULFID 28 631 INTERCHAIN (BY SIMILARITY).
FT DISULFID 83 94 BY SIMILARITY.
FT DISULFID 107 125 BY SIMILARITY.
FT DISULFID 141 144 BY SIMILARITY.
FT DISULFID 205 217 BY SIMILARITY.
FT DISULFID 228 247 BY SIMILARITY.
FT DISULFID 263 266 BY SIMILARITY.
FT DISULFID 327 339 BY SIMILARITY.
FT DISULFID 350 369 BY SIMILARITY.
FT CARBOHYD 82 82 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 242 242 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT VARSPIC 401 432 VSPPIAREQERDAETEQGTHGHWLHEKO -> RLIRA
FT CEYGRLSKAGAEPAPEQAESSQVKQ (in isoform
FT LMW).
FT /FTId=VSP_001263.
FT Missing (in isoform LMW).
FT /FTId=VSP_001264.
FT SEQUENCE 661 AA; 73102 MW; 774460258D58796E CRC64;
Query Match 65.2%; Score 430; DB 1; Length 661;
Best Local Similarity 65.5%; Pred. No. 1.8e-33;
Matches 81; Conservative 11; Mismatches 31; Indels 0; Gaps 0;
QY 1 GKDFVQPTKICVGCPRDIPNSPELEBTLTHITKLAENNAFYFKIDNVKARQVQV 60

Db 252 GDDLVALPKPCGCRDIPVDSPELKEVLGHSIAQLNAENHPFYKIDTVKATSQV 311
QY 61 AGKXYFIDFVARETCKSKESNEELTESCTKLGSLDCNANVYVVPWEKKIYPTVNCOP 120
Db 312 AGTKYVIEFIARETKCKSKESNTELAEDCEIKHLSGSLDCNANVYVWPENKVVPTVKCOA 371
QY 121 LGM 123
Db 372 LDM 374
RESULT 7
KNG_RAT
ID KNG_RAT STANDARD; PRT; 639 AA.
AC P08934; P08933;
DT 01-NOV-1988 (Rel. 09, Created)
DT 01-NOV-1988 (Rel. 09, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Kininogen precursor [Contains: Bradykinin].
GN KNG.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORMS HMW AND LMW).
RX MEDLINE=87137443; PubMed=3029068;
RA Kitagawa H., Kitamura N., Hayashida H., Miyata T., Nakanishi S.;
RT "Differing expression patterns and evolution of the rat kininogen
RT gene family.";
RL J. Biol. Chem. 262:2190-2198(1987).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORM LMW).
RX MEDLINE=86008264; PubMed=2413018;
RA Furuto-Rato S., Matsumoto A., Kitamura N., Nakanishi S.;
RT "Primary structures of the mRNAs encoding the rat precursors for
RT bradykinin and T-kinin. Structural relationship of kininogens with
RT major acute phase protein and alpha 1-cysteine proteinase
RT inhibitor.";
RL J. Biol. Chem. 260:12054-12059(1985).
RN [3]
RP SEQUENCE OF 1-65 FROM N.A.
RC STRAIN=Buffalo;
RX MEDLINE=87250580; PubMed=2439509;
RA Fung W.-P., Schreiber G.;
RT "Structure and expression of the genes for major acute phase alpha 1-
RT protein (thioesterin) and kininogen in the rat.";
RL J. Biol. Chem. 262:9298-9308(1987).
RN [4]
RP SEQUENCE OF 1-41 FROM N.A.
RC STRAIN=Wistar; TISSUE=Liver;
RX MEDLINE=87137465; PubMed=3818598;
RA Kageyama R., Kitamura N., Ohkubo H., Nakanishi S.;
RT "Differing utilization of homologous transcription initiation sites
RT of rat K and T kininogen genes under inflammation condition.";
RL J. Biol. Chem. 262:2345-2351(1987).
RN -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC HMW-kininogen plays an important role in blood coagulation by
CC helping to position optimally prekallikrein and factor XI next to
CC factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
CC induced aggregation of thrombocytes; (4) the active peptide
CC bradykinin that is released from HMW-kininogen shows a variety of
CC physiological effects: (4A) influence in smooth muscle
CC contraction, (4B) induction of hypotension, (4C) matrilysin and
CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
CC mediator of inflammation and causes (4E1) increase in vascular
CC permeability, (4E2) stimulation of nociceptors (4E3) release of
CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
CC a cardioprotective effect (directly via bradykinin action,
CC indirectly via endothelium-derived relaxing factor action); (5)
CC LMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
CC kininogen is in contrast to HMW-kininogen not involved in blood

DR PROSITE; PS00287; CYSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Vasodilator; Multigene family;
 KW Thiol protease inhibitor; Bradykinin; Acute phase; Signal.
 FT SIGNAL 1 18
 FT CHAIN 19 430 KININOGEN, T-II.
 FT CHAIN 19 375 HEAVY CHAIN.
 FT PEPTIDE 376 386 T-KININ.
 FT CHAIN 387 430 LIGHT CHAIN.
 FT DOMAIN 19 135 CYSTATIN-LIKE 1.
 FT DOMAIN 136 257 CYSTATIN-LIKE 2.
 FT DOMAIN 258 375 CYSTATIN-LIKE 3.
 FT DOMAIN 28 404 INTERCHAIN (BY SIMILARITY).
 FT DISULFID 83 94 BY SIMILARITY.
 FT DISULFID 107 125 BY SIMILARITY.
 FT DISULFID 141 144 BY SIMILARITY.
 FT DISULFID 205 217 BY SIMILARITY.
 FT DISULFID 228 247 BY SIMILARITY.
 FT DISULFID 263 266 BY SIMILARITY.
 FT DISULFID 327 339 BY SIMILARITY.
 FT DISULFID 350 369 BY SIMILARITY.
 FT CARBOHYD 82 82 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 126 126 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 326 326 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 430 AA; 47524 MW; 43EDF02D1BF55076 CRC64;
 Query Match 62.08; Score 409; DB 1; Length 430;
 Best Local Similarity 61.8%; Pred. No. 1.1e-31;
 Matches 76; Conservative 15; Mismatches 32; Indels 0; Gaps 0;
 Qy 1 GKDFVQPTKICVCPDIPNPSLEELTHITIKLNAENNAFFPKIDNVKARVQVY 60
 Db 252 GDDLFLSLPKFCGCPKPNIPVDSPELKEALGHSLAQLNAQHNLFFPKIDTVKATSQVY 311
 Qy 61 AGKYFIDFVARETSCSNEELTESCEKLGQSLDCAEYVVPWEKIYPTVNCOP 120
 Db 312 AGTYVIEFTARETSCSKQTNTLTADCTCKHGLQSLNCANVMYRPNKVVPTVRCQA 371
 Qy 121 LGM 123
 Db 372 LDM 374
 RESULT 9
 ID_KNTL_RAT
 AC P01038, P04081; STANDARD; PRT; 430 AA.
 DT 01-NOV-1986 (Rel. 03, Created)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE T-kininogen I precursor (Major acute phase protein) (Alpha-1-MAP)
 DE (Thiostatin) (Contains: T-kinin).
 GN MAP1.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=86008264; PubMed=2413018;
 RA Furuto-Kato S., Matsumoto A., Kitamura N., Nakanishi S.;
 RT "Primary structures of the mRNAs encoding the rat precursors for
 RT bradykinin and T-kinin. Structural relationship of kininogens with
 RT major acute phase protein and alpha 1-cysteine proteinase
 RT inhibitor.";
 RN J. Biol. Chem. 260:12054-12059(1985).
 FN [2]
 RP SEQUENCE OF 5-430 FROM N.A., AND PARTIAL SEQUENCE.
 RX MEDLINE=86008266; PubMed=2413019;
 RA Anderson K.P., Heath E.C.;
 RT "The relationship between rat major acute phase protein and the
 RT kininogens.";
 J. Biol. Chem. 260:12065-12071(1985).
 [3]
 RL SEQUENCE OF 7-430 FROM N.A.
 RN MEDLINE=85127561; PubMed=2578992;
 RX Cole T., Inglis A.S., Roxburgh C.M., Howlett G.J., Schreiber G.;
 RA "Major acute phase alpha 1-protein of the rat is homologous to bovine
 RT kininogen and contains the sequence for bradykinin: its synthesis is
 RT regulated at the mRNA level.";
 RL FEBS Lett. 182:57-61(1985).
 [4]
 RN SEQUENCE OF 1-65 FROM N.A.
 RX MEDLINE=87250580; PubMed=2439509;
 RA Fung W.-P., Schreiber G.;
 RT "Structure and expression of the genes for major acute phase alpha 1-
 RT protein (thiostatin) and kininogen in the rat.";
 RL J. Biol. Chem. 262:9298-9308(1987).
 CC -!- FUNCTION: Kininogens are plasma glycoproteins with a number of
 CC functions: (1) as precursor of the active peptide bradykinin and
 CC effect smooth muscle contraction, induction of hypotension and
 CC increase of vascular permeability. (2) They play a role in blood
 CC coagulation by helping to position optimally prekallikrein and
 CC factor XI next to factor XII. (3) They are inhibitor of thiol
 CC proteases.
 CC -!- SUBCELLULAR LOCATION: Extracellular.
 CC -!- TISSUE SPECIFICITY: Plasma.
 CC -!- INDUCTION: In response to an inflammatory stimulant. T-kininogen
 CC II synthesis is induced and the plasma concentration of
 CC T-kininogen I is raised.
 CC -!- PTM: AS T-KININ IS PRECEDED BY A MET INSTEAD OF AN ARG OR LYS, IT
 CC IS NOT RELEASED FROM ITS PRECURSOR BY EITHER TISSUE OR PLASMA
 CC KALLIKREIN.
 CC -!- MISCELLANEOUS: Rats express four types of kininogens: the
 CC classical HMW and LMW kininogens produced by alternative splicing
 CC of the same gene, and two additional LMW-like kininogens: T-I and
 CC T-II.
 CC -!- SIMILARITY: Contains 3 cystatin-like domains.
 CC -!- CAUTION: In addition to the conflicts described in the feature
 CC table, Ref.2 sequence differs from that shown in positions 257,
 CC 262, 268, 269, 295, 314, 315, 331, 332 and 389. In all those
 CC positions the alternate amino acid is the one present in T-II
 CC kininogen.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; M11883; AAA1489.1; -;
 CC EMBL; M1661; AAA41570.1; -;
 CC EMBL; M16454; AAA41568.1; -;
 CC EMBL; X02289; CAA26162.1; ALT_SEQ.
 CC PIR; A01286; KGRIT1.
 CC PIR; A23897; A23897.
 CC PIR; A27115; A27115.
 CC GlycosuiteDB; P01048; -;
 CC InterPro; IPR000010; Cystatin.
 CC Pfam; PF00031; cystatin; 3.
 CC SMART; SM00043; CY; 3.
 CC PROSITE; PS00287; CYSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Vasodilator; Multigene family;
 KW Thiol protease inhibitor; Bradykinin; Acute phase; Signal.
 FT SIGNAL 1 18
 FT CHAIN 19 430 KININOGEN, T-I.
 FT CHAIN 19 375 HEAVY CHAIN.
 FT PEPTIDE 376 386 T-KININ.
 FT CHAIN 387 430 LIGHT CHAIN.
 FT DOMAIN 19 135 CYSTATIN-LIKE 1.
 FT DOMAIN 136 257 CYSTATIN-LIKE 2.
 FT DOMAIN 258 375 CYSTATIN-LIKE 3.
 FT DISULFID 28 404 INTERCHAIN (BY SIMILARITY).


```

FT DISULFID 83 94 BY SIMILARITY.
FT DISULFID 107 125 BY SIMILARITY.
FT DISULFID 141 144 BY SIMILARITY.
FT DISULFID 205 217 BY SIMILARITY.
FT DISULFID 228 247 BY SIMILARITY.
FT DISULFID 263 266 BY SIMILARITY.
FT DISULFID 327 339 BY SIMILARITY.
FT DISULFID 350 369 BY SIMILARITY.
FT CARBOHYD 82 92 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 126 126 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 326 326 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CONFLICT 26 28 LNC -> MDR (IN REF. 2).
FT CONFLICT 55 55 V -> L (IN REF. 2).
FT CONFLICT 61 61 E -> Y (IN REF. 1).
FT CONFLICT 83 83 C -> Y (IN REF. 3).
FT CONFLICT 166 166 S -> F (IN REF. 2 AND 3).
FT CONFLICT 179 181 REV -> TKI (IN REF. 2).
FT CONFLICT 193 193 N -> D (IN REF. 2).
FT CONFLICT 212 212 S -> F (IN REF. 2).
FT CONFLICT 214 214 R -> H (IN REF. 3).
FT CONFLICT 229 229 T -> R (IN REF. 2).
FT CONFLICT 233 233 H -> Y (IN REF. 2).
FT CONFLICT 257 257 E -> S (IN REF. 2).
FT CONFLICT 262 262 N -> K (IN REF. 2).
FT CONFLICT 264 264 R -> F (IN REF. 2).
FT CONFLICT 268 269 RE -> KN (IN REF. 2).
FT CONFLICT 295 295 I -> L (IN REF. 2).
FT CONFLICT 314 315 VI -> TK (IN REF. 2).
FT CONFLICT 331 332 SK -> TN (IN REF. 2).
FT CONFLICT 389 389 R -> Q (IN REF. 2).
FT CONFLICT 414 414 R -> G (IN REF. 2 AND 3).
FT CONFLICT 415 415 A -> L (IN REF. 2).
FT CONFLICT 420 421 DH -> ER (IN REF. 3).
FT CONFLICT 430 430 P -> S (IN REF. 1).
SQ SEQUENCE 430 AA; 47715 MW; FAEBB78FAF4723C3 CRC64;

Query Match 60.8%; Score 401; DB 1; Length 430;
Best Local Similarity 61.8%; Pred. No. 6.3e-31;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;

QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNNAENATFYFKIDNVKARVQV 60
Db 252 GDLLELPKNGCRGPREIPVDSPKELGALHQAQLNAQNHIFYFKIDTVKATSOV 311
QY 61 AGKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQ 120
Db 312 AGVIVVIEFIARETNCQSKQKLTADCTKHLGQSLNCNANVYMRPWENKVVPTVRQA 371
QY 121 LQM 123
Db 372 LDM 374

RESULT 10
CYTF MOUSE
ID CYTF MOUSE STANDARD; PRT; 144 AA.
AC O89098;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Cystatin F precursor (Leukocystatin) (Cystatin 7) (Cystatin-like
DE metastasis-associated protein) (CMAP).
GN CS77.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Euthera; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98298157; PubMed=9632704;
RA Halfon S., Ford J., Foster J., Dowling L., Lucian L., Sterling M.,

```

```

RA Xu Y., Weiss M., Ikeda M., Liggett D., Helms A., Caux C., Lebecque S.,
RA "Leukocystatin, a new class II cystatin expressed selectively by
RA hematopoietic cells.",
RL J. Biol. Chem. 273:16400-16408(1998).
CC -I- FUNCTION: Inhibits papain and cathepsin L but with affinities
CC lower than other cystatins. May play a role in immune regulation
CC through inhibition of a unique target in the hematopoietic system.
CC -I- SUBCELLULAR LOCATION: Secreted (Probable).
CC -I- SIMILARITY: Belongs to the cystatin family.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; AF031826; AAC40140.1; -.
DR EMBL; AF031825; AAC40139.1; -.
DR HSPSP; P01034; I996.
DR MGD; MGI:1298217; Cst7.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00287; CYSTATIN; FALSE NEG.
KW Thiol protease inhibitor; Glycoprotein; Signal.
FT SIGNAL 1 18 POTENTIAL.
FT CHAIN 19 144 CYSTATIN F.
FT ACT_SITE 36 36 REACTIVE SITE.
FT SITE 80 84 SECONDARY AREA OF CONTACT.
FT DISULFID 98 109 BY SIMILARITY.
FT DISULFID 123 143 BY SIMILARITY.
SQ SEQUENCE 144 AA; 16380 MW; B5937334C1B4A89C CRC64;

Query Match 26.5%; Score 175; DB 1; Length 144;
Best Local Similarity 34.4%; Pred. No. 6.8e-10;
Matches 42; Conservative 23; Mismatches 45; Indels 8; Gaps 4;

QY 2 KDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNNAENATFYFKIDNVKARVQV 61
Db 27 KDLI---SSVKGFPTETNPGVLKARHSEVENFNCNTDIFLKESHVSKALVQVVK 83
QY 62 GKXYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVVVPWEKKI-YPTVN 117
Db 84 GLKYLEVIGRTTCKRTKTHQL-DNCDPQTNPALKRTLYCYSEVAVIPLHSEFVPVLL 142
QY 118 CQ 119
Db 143 CQ 144

RESULT 11
CYTF HUMAN
ID CYTF HUMAN STANDARD; PRT; 145 AA.
AC O76056; Q9UED4;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Cystatin F precursor (Leukocystatin) (Cystatin 7) (Cystatin-like
DE metastasis-associated protein) (CMAP).
GN CS77.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98406133; PubMed=9733783;
RA Ni J., Fernandez M.A., Danielsson L., Chillakuru R.A., Zhang J.,
RA Grubb A., Su J., Gentz R., Abrahamson M.;
RA "Cystatin F is a glycosylated human low molecular weight cysteine

```



```

RT RT proteinase inhibitor.";
RL J. Biol. Chem. 273:24797-24804(1998).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=98298157; PubMed=9632704;
RA Halfon S., Ford J., Foster J., Dowling L., Lucian L., Sterling M.,
RA Xu Y., Weiss M., Ikeda M., Liggett D., Helms A., Caux C., Lebecque S.,
RA Hannum C., Menon S., McClanahan T., Gorman D., Zurawski G.;
RA "Leukocystatin, a new class II cystatin expressed selectively by
RT hematopoietic cells.";
RL J. Biol. Chem. 273:16400-16408(1998).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=20399571; PubMed=10945474;
RA Morita M., Arakawa H., Yoshiuchi N.;
RT "Human homologue of murine CMAP.";
RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A.
RX MEDLINE=20399571; PubMed=10945474;
RA Morita M., Hara Y., Tanai Y., Arakawa H., Nishimura S.;
RT "Genomic construct and mapping of the gene for CMAP
RT Leukocystatin/Cystatin F, CSTF) and identification of a proximal
RT novel gene, BSCV (C20orf3).";
RL Genomics 67:87-91(2000).
RN [5]
RP SEQUENCE FROM N.A.
RX MEDLINE=21638749; PubMed=11780052;
RA Deloukas P., Matthews L.H., Ashurst J., Burton J., Gilbert J.G.R.,
RA Jones M., Stavrides G., Almeida J.P., Babbage A.K., Baggeley C.L.,
RA Bailey J., Barlow K.F., Bates K.N., Beare D.M.,
RA Beasley O.P., Bird C.P., Blakey S.E., Bridgman A.M., Brown A.J.,
RA Buck D., Burrill W.D., Butler A.P., Carder C., Carter N.P.,
RA Chapman J.C., Clamp M., Clark G., Clark L.N., Clark S.V., Clee C.M.,
RA Clegg S., Cobley V.E., Collier R.E., Connor R.E., Corby N.R.,
RA Coulson A., Coville G.J., Deadman R., Dhami P.D., Dunn M.,
RA Ellington A.G., Frankland J.A., Fraser A., French L., Garner P.,
RA Grafham D.V., Griffiths C., Griffiths M.N.D., Gwilliam R., Hall R.E.,
RA Hammond S., Harley J.L., Heath P.D., Ho S., Holden J.L., Howden P.J.,
RA Huckle E., Hunt A.R., Hunt S.E., Jekosch K., Johnson C.M., Johnson D.,
RA Kay M.P., Kimberley A.M., King A., Knights A., Laird G.K., Lawlor S.,
RA Lehaeslahe M.H., Leversha M.A., Lloyd C., Lloyd D.M., Lovell J.D.,
RA Marsh V.L., Martin S.L., McConachie L.J., McLeay K., McMurray A.A.,
RA Milne S.A., Mistry D., Moore M.J.P., Mullikin J.C., Nickerson T.,
RA Oliver K., Parker A., Patel R., Pearce T.A.V., Peck A.I.,
RA Phillimore B.J.C.T., Prathalingam S.R., Plumb R.W., Ramsay H.,
RA Rice C.M., Ross M.T., Scott C.E., Sehra H.K., Showkneen R., Sims S.,
RA Skuce C.D., Smith M.L., Soderlund C., Steward C.A., Sulston J.E.,
RA Swann R.M., Sycamore N., Taylor R., Tee L., Thomas D.W., Thorpe A.,
RA Tracey A., Tromans A.C., Vaudin M., Wall M., Wallis J.M.,
RA Whitehead S.L., Whittaker P., Willey D.L., Williams L., Williams S.A.,
RA Wilming L., Wray P.W., Hubbard T., Durbin R.M., Bentley D.R., Beck S.,
RA Rogers J.;
RT "The DNA sequence and comparative analysis of human chromosome 20.";
RL Nature 414:865-871(2001).
RN [6]
RP SEQUENCE FROM N.A.
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Udwin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaby S.J.,
RA Boek S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettner M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,

```

```

RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -1- FUNCTION: Inhibits papain and cathepsin L but with affinities
CC lower than other cystatins. May play a role in immune regulation
CC through inhibition of a unique target in the hematopoietic system.
CC -1- SUBCELLULAR LOCATION: Secreted (Probable).
CC -1- TISSUE SPECIFICITY: Primarily expressed in peripheral blood cells
CC and spleen.
CC -1- SIMILARITY: Belongs to the cystatin family.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AF036342; AAC35747.1; -.
CC EMBL; AF031824; AAC39788.1; -.
CC EMBL; AB015225; BAA34941.1; ALT_INIT.
CC EMBL; AB029636; BAB11886.1; ALT_INIT.
CC EMBL; AL035661; CAB75498.1; -.
CC EMBL; BC015507; AAI15507.1; ALT_INIT.
CC HSP; F01034; IG96.
CC MIM; 603253; -.
CC GO; GO:0004869; F:cysteine protease inhibitor activity; TAS.
CC GO; GO:0006955; P:immune response; TAS.
CC InterPro; IPR000010; Cystatin.
CC Pfam; PF00031; cystatin; 1.
CC SMART; SM00043; C1; 1.
CC GO; GO:0004869; F:cysteine protease inhibitor activity; TAS.
CC PROSITE; PS00287; CYSTATIN; FALSE_NEG.
CC Thiol protease inhibitor; Glycoprotein; Signal.
CC SIGNAL 1 19 POTENTIAL.
CC CHAIN 20 145 CYSTATIN F.
CC ACT SITE 37 37 REACTIVE SITE.
CC SITE 81 85 SECONDARY AREA OF CONTACT.
CC DISULFID 99 110 BY SIMILARITY.
CC DISULFID 124 144 BY SIMILARITY.
CC CARBOHYD 62 62 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC CARBOHYD 115 115 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC SQ SEQUENCE 145 AA; 16454 MW; B2BCC4F76857CB0F CRC64;
Query Match 25.1%; Score 165.5; DB 1; Length 145;
Best Local Similarity 32.5%; Pred. No. 5.5e-09;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;
QY 9 TKICVCGPRDIPNPSLEBETLTHITKLNAENNAFFKIDNVKARVQVWAGKYFID 68
Db 32 SRVKPGFKTIKNDPGLQARYSVEKFNCTNDMLFKESRITRALQIVKGLKYLE 91
QY 69 FVARETTCESNEELTESCE---TKKLGQSLDCNAEVVVPWEKKI-VPTVNC 118
Db 92 VEIGRTTCKKNQHLRL-DDCDFQTNHTLKTLSYSEVWVWVWLFQHFVEVLR 144
RESULT 12
CYTM HUMAN
ID CYTM HUMAN STANDARD; PRT; 149 AA.
AC Q15828;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Cystatin M precursor (Cystatin E).
GN CST6.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;

```

RN SEQUENCE FROM N.A.
 RP MEDLINE=97150844; PubMed=8995380;
 RX Sotiropoulos G., Anisowicz A., Sager R.;
 RA "Identification, cloning, and characterization of cystatin M, a novel
 RT cysteine proteinase inhibitor, down-regulated in breast cancer.";
 RL J. Biol. Chem. 272:903-910(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RP MEDLINE=97256812; PubMed=9099741;
 RA Ni J., Abrahamson M., Zhang M., Fernandez M.A., Grubb A., Su J.,
 RA Yu G.L., Li Y., Parmelee D., Xing L., Coleman T.A., Gentz S.,
 RA Thakura R., Nguyen N., Hesselberg M., Gentz R.;
 RT "Cystatin E is a novel human cysteine proteinase inhibitor with
 RT structural resemblance to family 2 cystatins.";
 RL J. Biol. Chem. 272:10853-10858(1997).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Prostate;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diachenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavert T.L., Scheetz T.E.,
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Frange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahey J., Helton E., Kettelman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzyzinski M.I., Skalska U., Smalhus D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [4]
 RP CHARACTERIZATION, AND TISSUE SPECIFICITY.
 RX MEDLINE=21246880; PubMed=11348457;
 RA Zeeuwen P.L., Van Vlijmen-Willems I.M., Jansen B.J., Sotiropoulos G.,
 RA Curfs J.H., Meis J.F., Janssen J.J., Van Ruissen P., Schalkwijk J.;
 RT "Cystatin M/E expression is restricted to differentiated epidermal
 RT keratinocytes and sweat glands: a new skin-specific proteinase
 RT inhibitor that is a target for cross-linking by transglutaminase.";
 RL J. Invest. Dermatol. 116:693-701(2001).
 CC -1- FUNCTION: Shows moderate inhibition of cathepsin B but is not
 CC active against cathepsin C.
 CC -1- SUBCELLULAR LOCATION: Secreted.
 CC -1- TISSUE SPECIFICITY: Restricted to the stratum granulosum of normal
 CC skin, the stratum granulosum/spinosum of psoriatic skin, and the
 CC secretory coils of eccrine sweat glands. Low expression levels are
 CC found in the nasal cavity.
 CC -1- PTM: Substrate for transglutaminases. Acts as an acyl acceptor but
 CC not as an acyl donor.
 CC -1- SIMILARITY: Belongs to the cystatin family.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC
 DR EMBL; U62800; AAB06566.1; -
 DR EMBL; U81233; AAB61305.1; -
 DR EMBL; BC031334; AAB31334.1; -
 DR HSSP; P01038; 1CEW.
 DR Genew; HGNC:2478; CST6.

DR MIN; 601891; -
 DR GO; GO:0004869; F:cysteine protease inhibitor activity; TAS.
 DR GO; GO:0007345; P:embryogenesis and morphogenesis; TAS.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; Cystatin; 1.
 DR SMART; SM00343; CY; 1.
 DR PROSITE; PS00287; CYSTATIN; 1.
 KW Thiol protease inhibitor; Signal; Glycoprotein.
 FT SIGNAL 1 28 PROBABLE.
 FT CHAIN 29 149 CYSTATIN M.
 FT ACT SITE 36 36 REACTIVE SITE.
 FT SITE 80 84 SECONDARY AREA OF CONTACT.
 FT DISULFID 98 113 BY SIMILARITY.
 FT DISULFID 126 146 BY SIMILARITY.
 FT CARBOHYD 137 137 N-LINKED (GLCNAC...) (POTENTIAL).
 SQ SEQUENCE 149 AA; 2076A78BFC9FAC8C CRC64;
 Query Match 21.0%; Score 138.5; DB 1; Length 149;
 Best Local Similarity 31.5%; Pred. No. 21e-06;
 Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
 Qy 8 PTKICVCPRIDIPNPSPELEETLTHITIKLNENNAFYFKIDNVKARVQVVGKKYFI 67
 Db 30 PQSRVWGLRDLSPDPQVQKAAQAAVSYNMGNSIYYFRDTHITIKAQSLVAGIKYFI 89
 Qy 68 DFVARETTCSKE---SNEELTESCETKLGQ--SLDCNAEVVVPWE 109
 Db 90 TMENGSTDCRKTVTGCHVDLI--TCPAAGAQQEKLRCDFEVLVVPWQ 136
 RESULT 13
 CYTC_BOVIN STANDARD; PRT; 148 AA.
 ID CYTC_BOVIN STANDARD; PRT; 148 AA.
 AC P01035;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 15-JUL-1999 (Rel. 38, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Cystatin C precursor (Colostrum thiol proteinase inhibitor).
 GN CST3.
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.; SEQUENCE OF 66-83, AND CHARACTERIZATION.
 RC TISSUE=Cerebrospinal fluid, and Choroid plexus;
 RX MEDLINE=98094199; PubMed=9434110;
 RA Olsson S.-L., Ek B., Wilm M., Broberg S., Rask L., Bjoerk I.;
 RT "Molecular cloning and N-terminal analysis of bovine cystatin C
 RT identification of a full-length N-terminal region.";
 RL Biochim. Biophys. Acta 1343:203-210(1997).
 RN [2]
 RP SEQUENCE OF 37-148
 RX MEDLINE=8521205; PubMed=3891407;
 RA Hirado M., Tsunawawa S., Sakiyama F., Niinobe M., Fujii S.;
 RT "Complete amino acid sequence of bovine colostrum low-Mr cysteine
 RT proteinase inhibitor.";
 RL FEBS Lett. 186:41-45(1985).
 CC -1- FUNCTION: This is a thiol proteinase inhibitor.
 CC -1- MASS SPECTROMETRY: MW=13420, METHOD=VALDI.
 CC -1- SIMILARITY: Belongs to the cystatin family.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC
 DR EMBL; Y10811; CAA71771.1; -
 DR HSSP; P01034; 1C96.

```
DR InterPro; IPR000010; Cystatin.  
DR Pfam; PF000031; cystatin; 1.  
DR SMART; SM00043; CY; 1.  
DR PROSITE; PS00287; CYSTATIN; 1.  
KW Thiol protease inhibitor; Signal; Pyrrolidone carboxylic acid.  
FT SIGNAL 1 30 PROBABLE.  
FT CHAIN 31 148 CYSTATIN C.  
FT MOD_RES 31 31 PYRROLIDONE CARBOXYLIC ACID (PROBABLE).  
FT ACT_SITE 40 40 REACTIVE SITE.  
FT SITE 84 88 SECONDARY AREA OF CONTACT.  
FT DISULFID 102 112 BY SIMILARITY.  
FT DISULFID 126 146 BY SIMILARITY.  
SQ SEQUENCE 148 AA; 16265 MW; EE740FE37CFB9F0E CRC64;  
  
Query Match 20.6%; Score 136; DB 1; Length 148;  
Best Local Similarity 30.6%; Pred. No. 3.7e-06;  
Matches 34; Conservative 22; Mismatches 31; Indels 24; Gaps 5;  
  
QY 22 NSPELEETLTHITKLNAENNAFFKIDNVKARQVQVAGKKYFIDFVARETTCSKESN 81  
DB 48 NEEGVQEAISFAVSEFNKRSNDAYOSRVVRVRAKQVVGNGVFLDVELGRITCTK--S 105  
QY 82 BELTESC-----ETKLGQSLDCNAEVYVWPWEKKIYPTVN-----CQ 119  
DB 106 QANLDSCTPHNPKREKL-----CSFQVYVWPWN-----TINLVKSCQ 147  
  
RESULT 14  
CYTC_MACMU  
ID CYTC_MACMU STANDARD; PRT; 146 AA.  
AC Q19032;  
DT 15-JUL-1998 (Rel. 36, Created)  
DT 15-JUL-1998 (Rel. 36, Last sequence update)  
DT 28-FEB-2003 (Rel. 41, Last annotation update)  
DE Cystatin C precursor.  
GN CST3.  
OS Macaca mulatta (Rhesus macaque).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;  
OC Cercopithecoidea; Macaca.  
OX NCBI_TaxID=9544;  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=97054523; PubMed=8898820;  
RA Wei L.H., Walker L.C., Levy E.;  
RT "Cystatin C. Icelandic-like mutation in an animal model of  
RT cerebrovascular beta-amyloidosis.";  
RL Stroke 27:2080-2085(1996).  
CC -!- FUNCTION: As an inhibitor of cysteine proteinases, this protein is  
CC thought to serve an important physiological role as a local  
CC regulator of this enzyme activity.  
CC -!- SIMILARITY: Belongs to the cystatin family.  
CC  
CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration  
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -  
CC the European Bioinformatics Institute. There are no restrictions on its  
CC use by non-profit institutions as long as its content is in no way  
CC modified and this statement is not removed. Usage by and for commercial  
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/  
CC or send an email to license@isb-sib.ch).  
CC  
CC EMBL; U52028; AAB64051.1; -.  
CC HSP; P01034; I936.  
CC InterPro; IPR000010; Cystatin.  
CC Pfam; PF00031; cystatin; 1.  
CC SMART; SM00043; CY; 1.  
CC PROSITE; PS00287; CYSTATIN; 1.  
KW Thiol protease inhibitor; Amyloid; Signal.  
FT SIGNAL 1 26 BY SIMILARITY.  
FT CHAIN 27 146 CYSTATIN C.  
FT ACT_SITE 37 37 REACTIVE SITE.  
FT SITE 81 85 SECONDARY AREA OF CONTACT.  
FT DISULFID 99 109 BY SIMILARITY.  
FT DISULFID 123 143 BY SIMILARITY.  
SQ SEQUENCE 146 AA; 15946 MW; 08196353C0306AA3 CRC64;  
  
Query Match 20.1%; Score 132.5; DB 1; Length 146;  
Best Local Similarity 29.0%; Pred. No. 7.8e-06;  
Matches 29; Conservative 21; Mismatches 45; Indels 5; Gaps 2;  
  
QY 13 VGCPRDIPNTSPLEETLTHITKLNAENNAFFKIDNVKARQVQVAGKKYFIDFVAR 72  
DB 36 LGGEMDASVEEGVRRALDFAVSEYNKASNDMVHSSALQVWRARKQIVAGVNYFLDVMG 95  
QY 73 ETTCSKESNEELTESC-----ETKLGQSLDCNAEVYVWPWE 109
```

Page 14

Search completed: September 24, 2004, 14:09:12
Job time : 23.364 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:05:18 ; Search time 35.424 Seconds
(without alignments)
1095.549 Million cell updates/sec

Title: US-10-661-784-1

Perfect score: 660

Sequence: 1 GKDFVQPTKICVGCPRDIP.....YVVPWEKKIYTVNCQPLGM 123

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL 25:**

- 1: sp_archaea:**
- 2: sp_bacteria:**
- 3: sp_fungi:**
- 4: sp_human:**
- 5: sp_invertebrate:**
- 6: sp_mammal:**
- 7: sp_mhc:**
- 8: sp_organelle:**
- 9: sp_phase:**
- 10: sp_plant:**
- 11: sp_podent:**
- 12: sp_virus:**
- 13: sp_vertebrate:**
- 14: sp_unclassified:**
- 15: sp_virus:**
- 16: sp_bacteriap:**
- 17: sp_archaeap:**

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	404	61.2	140	6 Q7YRP6	Q7YRP6 sus scrofa
2	402	60.9	423	11 P70517	P70517 rattus norv
3	399	60.5	430	11 Q63581	Q63581 rattus norv
4	175	26.5	167	11 Q9QWL5	Q9QWL5 mus musculu
5	165.5	25.1	167	4 Q7Z4J8	Q7Z4J8 homo sapien
6	152.5	23.1	462	13 Q7Z191	Q7Z191 xenopus lae
7	152.5	23.1	462	13 Q7SVH2	Q7SVH2 xenopus lae
8	152.5	23.1	465	13 Q801B5	Q801B5 xenopus lae
9	129.5	19.6	140	11 Q9EPX9	Q9EPX9 mus musculu
10	122.5	18.6	455	13 Q80058	Q80058 brachydanio
11	118	17.9	464	13 Q801Z5	Q801Z5 cyprinus ca
12	115.5	17.5	388	11 Q8CB17	Q8CB17 mus musculu
13	113.5	17.2	148	5 Q9NH95	Q9NH95 litomosoid
14	113	17.1	140	11 Q80Y72	Q80Y72 mus musculu
15	111	16.8	146	11 Q8K397	Q8K397 mus musculu
16	111	16.8	149	11 Q9D1B1	Q9D1B1 mus musculu

ALIGNMENTS

RESULT 1

Q7YRP6 ID Q7YRP6 PRELIMINARY; PRT; 140 AA.
AC Q7YRP6;
DT 01-OCT-2003 (TREMREL. 25, Created)
DT 01-OCT-2003 (TREMREL. 25, Last sequence update)
DT 01-OCT-2003 (TREMREL. 25, Last annotation update)
DE Low molecular weight kininogen (Fragment).
GN KNG.
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
OX NCBI_TaxID=9823;
RN [1]
RP SEQUENCE FROM N.A.
RA Vonnahme K.A., Fernando S.C., Ross J.A., Ashworth M.D., DeSilva U.,
RA Malayer J.R., Geisert R.D.;
RT "Porcine Endometrial and Conceptus Expression of Kininogens and Plasma
RT Kallikrein in Cyclic and Pregnant Gilts."
RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY213363; AAP85260.1; -;
FT NON_TER 1
FT NON_TER 140
SQ SEQUENCE 140 AA; 15650 MW; 177837836603F777 CRC64;

Query Match 61.2%; Score 404; DB 6; Length 140;

Best Local Similarity 78.2%; Pred. No. 5.4e-33;

Matches 79; Conservative 5; Mismatches 17; Indels 0; Gaps 0;

Qy 22 NSPELETHTHTITKLNAENNATFYFKIDNVKARVQVAGKKYFDVFARETTCSKSN 81

Db 1 DSPDSEPLNHSIAKLNAENNAVFYFKIGPVEKATVQVAGKKYSIVFARETTCSKSN 60

Qy 82 ELTSCETCKKLGOSLDCAEYVYVVPWEKKIYTVNCQPLG 122

Db 61 ELTSCETCKKQQLKCNASVYVVPWEKKIYTVNCQPLG 101

RESULT 2

```

P70517
ID P70517 PRELIMINARY; PRT; 423 AA.
AC MEDLINE=85149311; PubMed=2579644;
DT 01-FEB-1997 (TREMBlrel. 02, Created)
DT 01-FEB-1997 (TREMBlrel. 02, Last sequence update)
DE Major acute phase alpha(1)-protein in the rat: Structure, molecular
DE Major acute phase alpha-1 protein precursor (Fragment).
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RA Cole T.;
RL Submitted (OCT-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=85149311; PubMed=2579644;
RA Cole T., Inglis A., Nagashima M., Schreiber G.;
RT "Major acute-phase alpha(1)-protein in the rat: Structure, molecular
RT cloning, and regulation of mRNA levels.";
RL Biochem. Biophys. Res. Commun. 126:719-724(1985).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=85127561; PubMed=2578992;
RA Cole T., Inglis A.S., Roxburgh C.M., Howlett G.J., Schreiber G.;
RT "Major acute phase alpha1-protein of the rat is homologous to bovine
RT kininogen and contains the sequence for bradykinin: its synthesis is
RT regulated at the mRNA level.";
RL FEBS Lett. 182:57-61(1985).
DR EMBL; K02814; AAA41569.1; -.
DR PIR; S68034; S68034.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 3.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Signal.
FT NON TER 1 1
FT SIGNAL <1 11 POTENTIAL.
FT CHAIN 12 423 POTENTIAL.
FT CHAIN 371 379 POTENTIAL.
SQ SEQUENCE 423 AA; 46905 MW; F958BD3198547949 CRC64;

Query Match 60.9%; Score 402; DB 11; Length 423;
Best Local Similarity 61.8%; Pred. No. 3.1e-32;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;

QY 1 GKDFVQPPKICVGCPRDIPNTSPLEETLTHITKLNAENNATFYFKIDNVKKARVQV 60
DQ 245 GDDLPELLPNCRGCPREIPVDSPELKEALGHSIARLNAQHNIIFYFKIDTVKATSOV 304
QY 61 AGKXYFIDFVARETTCKESNEBELTESCTKKLGSLDCNAEYVVPWEKKIYPTVNCOP 120
DQ 305 AGVIYVIEFIARETNGSKQKTELTADCTCKHLGSLNCNANVYMRPWENKVPVRCQA 364
QY 121 LGM 123
DQ 365 LDM 367

RESULT 3
ID Q63581 PRELIMINARY; PRT; 430 AA.
AC Q63581;
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
DE Rat T-kininogen (T-KG).
DE Rat T-kininogen (T-KG).
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

```

```

OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90034172; PubMed=2806908;
RA Anderson K.P., Croyle M.L., Lingrel J.B.;
RT "Primary structure of a gene encoding rat T-kininogen.";
RL Gene 81:119-128(1989).
DR EMBL; M29090; AAA42251.1; JOINED.
DR EMBL; M29083; AAA42251.1; JOINED.
DR EMBL; M29084; AAA42251.1; JOINED.
DR EMBL; M29091; AAA42251.1; JOINED.
DR EMBL; M29085; AAA42251.1; JOINED.
DR EMBL; M29086; AAA42251.1; JOINED.
DR EMBL; M29087; AAA42251.1; JOINED.
DR EMBL; M29088; AAA42251.1; JOINED.
DR EMBL; M29089; AAA42251.1; JOINED.
DR PIR; S68034; S68034.
DR PIR; S68035; S68035.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 3.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
SQ SEQUENCE 430 AA; 47618 MW; 45508DEF4BDC978C CRC64;

Query Match 60.5%; Score 399; DB 11; Length 430;
Best Local Similarity 61.8%; Pred. No. 6.3e-32;
Matches 76; Conservative 13; Mismatches 34; Indels 0; Gaps 0;

QY 1 GKDFVQPPKICVGCPRDIPNTSPLEETLTHITKLNAENNATFYFKIDNVKKARVQV 60
DQ 252 GDDLPELLPNCRGCPREIPVDSPELKEALGHSIARLNAQHNIIFYFKIDTVKATSOV 311
QY 61 AGKXYFIDFVARETTCKESNEBELTESCTKKLGSLDCNAEYVVPWEKKIYPTVNCOP 120
DQ 312 AGVIYVIEFIARETNGSKQKTELTADCTCKHLGSLNCNANVYMRPWENKVPVRCQA 371
QY 121 LGM 123
DQ 372 LDM 374

RESULT 4
ID Q9QWL5 PRELIMINARY; PRT; 167 AA.
AC Q9QWL5;
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)
DE Murine CMAP (CYSTATIN F) (LEUKOCYSTATIN).
DE MURINE CMAP OR CST7.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA Morita M., Arakawa H., Yoshiuchi N.;
RT "A novel cystatin-like metastasis associated gene.";
RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX STRAIN=C57BL/6J; TISSUE=Embryo;
RX MEDLINE=21085660; PubMed=11217851;
RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
RA Azawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamana I.,
RA Saito T., Okazaki Y., Gojibori T., Bono H., Kasukawa T., Saito R.,
RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant I.,
RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
RA Kiehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
RA Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,

```

```

RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,
RA Hayashizaki Y.,
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:685-690(2001).
DR EMBL; AB015224; BAA34940.1; -.
DR EMBL; AK004420; BAB23298.1; -.
DR HSSP; P01034; 1G96.
DR MGD; MGI:1298217; Cst7.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; Cyt; 1.
SQ SEQUENCE 167 AA; 18847 MW; 61F776D8445095FE CRC64;
Query Match 26.5%; Score 175; DB 11; Length 167;
Best Local Similarity 34.4%; Pred. No. 8.3e-10;
Matches 42; Conservative 23; Mismatches 49; Indels 8; Gaps 4;
Qy 2 KDFVQPTKICVGPRIPTNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVVA 61
Db 50 KDLI---SSVKGPFKTIETNPGVLKAARSHVEKFNNTDIFLFKESHVSKALVQVVK 106
Qy 62 GKYPIDFVARETTCSKESNEELTESC---TKLQSLDCNAEVVVPWEKTI-YPTVN 117
Db 107 GLKTNLEVKIGRTTCRKMHQL-DNCDFTQNPALKRTLYCYSEVWPVLHSEFVPLL 165
Qy 118 CQ 119
Db 166 CQ 167

RESULT 5
Q7Z4J8 PRELIMINARY; PRT; 167 AA.
AC Q7Z4J8;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cystatin F (Leukocystatin).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Ventral midgut;
RA Kainine N., Chen X., Rolfs A., Halleck A., Hines L., Eisenstein S.,
RA Koundinya M., Raphael J., Moreira D., Kelley T., LaBaer J., Lin Y.,
RA Phelan M., Farmer A.;
RL Submitted (AUG-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BT009825; AAP88827.1; -.
SQ SEQUENCE 167 AA; 18857 MW; E339025A5BD60177 CRC64;
Query Match 25.1%; Score 165.5; DB 4; Length 167;
Best Local Similarity 32.5%; Pred. No. 7.5e-09;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;
Qy 9 TKICVGPRIPTNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVQVAGKYFID 68
Db 54 SRVKGPFKTIKTNDPGVQLQARYSVKFNNTDMLFKESRITRALVQIVKGLKYLE 113
Qy 69 FVARETTCSKESNEELTESC---TKLQSLDCNAEVVVPWEKTI-YPTVNC 118
Db 114 VEIGRTICKNQHLRL-DDCDFQTNHTILKQTLSCYSEVWPVLQHFVFLRC 166

RESULT 6
Q7ZV91 PRELIMINARY; PRT; 462 AA.
AC Q7ZV91;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Similar to fetuin B.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Embryo;
RA Klein S., Strausberg R.;
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC043891; AAH43891.1; -.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; Cyt; 2.
SQ SEQUENCE 462 AA; 53185 MW; D7BAD339961739FB CRC64;
Query Match 23.1%; Score 152.5; DB 13; Length 462;
Best Local Similarity 38.8%; Pred. No. 5e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;
Qy 8 PTKICVGPRIPTNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVQVAGK 63
Db 142 PGVILSTCP-DCFTANEETPTITADTLIAEYKNSNNTFYFKIDHIERVRSQWVGP 200
Qy 64 KYFIDFVARETTCSKESNEELTESC 88
Db 201 SYFIQTIKETDCMKTQENVVLSNC 225

RESULT 7
Q7SYH2 PRELIMINARY; PRT; 462 AA.
AC Q7SYH2;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cystatin domain fetuin-like protein.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Ventral midgut;
RA Costa R.M.B., Mason J., Lee M., Amaya E., Zorn A.M.;
RA "Novel gene expression domains reveal early patterning of the Xenopus
RT endoderm.";
RL Submitted (MAR-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY260732; AAP82289.1; -.
SQ SEQUENCE 462 AA; 53186 MW; 796F92774CC27721 CRC64;
Query Match 23.1%; Score 152.5; DB 13; Length 462;
Best Local Similarity 38.8%; Pred. No. 5e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;
Qy 8 PTKICVGPRIPTNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVQVAGK 63
Db 142 PGVILSTCP-DCFTANEETPTITADTLIAEYKNSNNTFYFKIDHIERVRSQWVGP 200
Qy 64 KYFIDFVARETTCSKESNEELTESC 88
Db 201 SYFIQTIKETDCMKTQENVVLSNC 225

```

```

RESULT 8
Q801E5 PRELIMINARY; PRT; 465 AA.
AC Q801E5;
DT 01-JUN-2003 (TREMELrel. 24, Created)
DT 01-JUN-2003 (TREMELrel. 24, Last sequence update)
DT 01-OCT-2003 (TREMELrel. 25, Last annotation update)
DE Hypothetical histidine-rich protein (Fragment).
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22480013; PubMed=12591597;
RA Chen Y., Jurgens K., Hollemann T., Clausen M., Ramadori G.,
RA Pieler T.;
RT "Cell-autonomous and signal-dependent expression of liver and
RT intestine marker genes in pluripotent precursor cells from Xenopus
RT embryos."; 120:277-288 (2003).
RL Mech. Dev. 120:277-288 (2003).
DR EMBL; AY18284; AAC31610.1; -.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; CY; 2.
KW Hypothetical protein.
FT NON_TER
SQ SEQUENCE 465 AA; 53528 MW; 0B403AB4F78BFD4 CRC64;

Query Match 23.1%; Score 152.5; DB 13; Length 465;
Best Local Similarity 38.8%; Pred. No. 5e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

QY 8 PTKVCGPRDIPINSPELEETLTWT---ITKLAENNAATFYKIDNVKARQVQVAGK 63
DB 145 PGVILSTCP-DCPTANEETITPTITAEITLAEVKNKSNNTFYKIDHIERVRSQWVGP 203
QY 64 KYFIDFVARETTCSKESNEELTESC 88
DB 204 SYFIQFTIKETDCKTQENVLSNC 228

RESULT 9
Q9EPX9 PRELIMINARY; PRT; 140 AA.
AC Q9EPX9;
DT 01-MAR-2001 (TREMELrel. 16, Created)
DT 01-MAR-2001 (TREMELrel. 16, Last sequence update)
DT 01-JUN-2003 (TREMELrel. 24, Last annotation update)
DE Cystatin C.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=BALB/c;
RX MEDLINE=21010502; PubMed=11144350;
RA Taupin P.J., Ray J., Fischer W.H., Suhr S.T., Hakansson K., Grubb A.,
RA Gage F.H.;
RT "FGF-2-Responsive neural stem cell proliferation requires CCG, a novel
RT autocorine/paracrine cofactor.";
RL Neuron 28:385-397 (2000).
DR EMBL; AF311741; AAG40263.1; -.
DR HSSP; P01034; 1G96.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00287; CYSTATIN; 1.
FT CHAIN 21 140 CYSTATIN C.

```

```

FT VARIANT 16 16 A -> G.
FT VARIANT 84 84 L -> P.
SQ SEQUENCE 140 AA; 15517 MW; 3A563406DD58D785 CRC64;

Query Match 19.6%; Score 129.5; DB 11; Length 140;
Best Local Similarity 30.0%; Pred. No. 2.6e-05;
Matches 30; Conservative 21; Mismatches 44; Indels 5; Gaps 3;

QY 13 VCGPRDIPINSPELEETLTHTITKLAENNAATFYKIDNVKARQVQVAGKVFIDFVAR 72
DB 30 LGAPEADANEGRVRRALDFAVSEYKNGSDAHSGRAIQVVRKQOLVAGVNFELDEVG 89
QY 73 ETTCSKESNEELTESC---ETKKLGSLDCNAEAVVVPWE 109
DB 90 RYTCTK-SQTNLTD-CFFHDQPHLMKALCSFQIYVVPWK 127

RESULT 10
Q800S8 PRELIMINARY; PRT; 455 AA.
AC Q800S8;
DT 01-JUN-2003 (TREMELrel. 24, Created)
DT 01-JUN-2003 (TREMELrel. 24, Last sequence update)
DT 01-OCT-2003 (TREMELrel. 25, Last annotation update)
DE Fetuin-A.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RA Jia F.;
RT "Danio rerio fetuin-A.";
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY217758; AAC64483.1; -.
DR GO; GO:0005874; C:microtubule; IEA.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR GO; GO:0005198; F:structural molecule activity; IEA.
DR GO; GO:0007018; P:microtubule-based movement; IEA.
DR InterPro; IPR002453; Beta tubulin.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00228; TUBULIN B AUTOREG; 1.
SQ SEQUENCE 455 AA; 50627 MW; D822872926BA2ACB CRC64;

Query Match 18.6%; Score 122.5; DB 13; Length 455;
Best Local Similarity 26.4%; Pred. No. 0.00052;
Matches 29; Conservative 22; Mismatches 44; Indels 15; Gaps 3;

QY 10 KICVCGPRDIPINSPELEETLTHTITKLAENNAATFYKIDNVKARQVQVAGKVFID 68
DB 140 KKCPDCPGLPLPHEPKALESVNALAKFNKQSNHKSFYKLMVEVGRISQWMPMGOSYFTQ 199
QY 69 FVARETTCSKESNEELTES-----CETKKLG-QSLDCNAEAVV 104
DB 200 FAIMETNCTKKDAPQNPEACKALCGDQATYGFCKSKVGSEPEVECEIY 249

RESULT 11
Q801Z5 PRELIMINARY; PRT; 464 AA.
AC Q801Z5;
DT 01-JUN-2003 (TREMELrel. 24, Created)
DT 01-JUN-2003 (TREMELrel. 24, Last sequence update)
DT 01-OCT-2003 (TREMELrel. 25, Last annotation update)
DE Fetuin long form.
OS Cyprinus carpio (Common carp).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Cyprinus.

```



```

OX NCBI_TaxID=7962;
RN [1]
RP SEQUENCE FROM N.A.
RA Teai P.-L., Chang G.-D., Huang C.-J.;
RT "Purification and cloning of carp fetuin.";
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY225965; AA074862.1; -.
DR GO; GO:0005874; C:microtubule; IEA.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR GO; GO:0005198; F:structural molecule activity; IEA.
DR GO; GO:0007018; P:microtubule-based movement; IEA.
DR InterPro; IPR002453; Beta tubulin.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00228; TUBULIN_B_AUTOREG; 1.
SQ SEQUENCE 464 AA; 51698 MW; 7A54F71B44050895 CRC64;

Query Match 17.9%; Score 118; DB 13; Length 464;
Best Local Similarity 24.2%; Pred. No. 0.0015;
Matches 31; Conservative 27; Mismatches 50; Indels 20; Gaps 5;

QY 12 CVGCPRIPTNSPELEETLTHITKLNAENNATFYFKIDNVKARVQ-VVAGKKYFIDFV 70
DB 142 CPDCEGLPLHDPKGLSVKTLQKFNKESDHKSYPKLMVEVGRISTQWPFSGQSFTSQPA 201
QY 71 ARETTCSE---SNEELTES-----CETKLG-QSLDCNVAEYVV-----PREKK 111
DB 202 IMETNCTNKEAPQNEESKALCGEKARYGFKSTKVGIEPEVECEIYEAKNITHPMKHP 261
QY 112 IYPTVNCQ 119
DB 262 AQSRDCK 269

RESULT 12
Q8CB17 PRELIMINARY; PRT; 388 AA.
AC Q8CB17
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DR Fetuin beta.
GN FETUB.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=C57BL/6J; TISSUE=Vagina;
RX MEDLINE=22354683; PubMed=12466851;
RA The FANTOM Consortium,
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs.";
RL Nature 420:563-573(2002).
DR EMBL; AK037043; BAC29682.1; -.
DR MGD; MGI:1690221; Fetub.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR InterPro; IPR001363; Fetuin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; CY; 2.
DR PROSITE; PS01254; FETUIN 1; 1.
DR PROSITE; PS01255; FETUIN 2; 1.
SQ SEQUENCE 388 AA; 42742 MW; 78CFAD73A8D8DC22 CRC64;

Query Match 17.5%; Score 115.5; DB 11; Length 388;
Best Local Similarity 26.7%; Pred. NO. 0.0022;
Matches 31; Conservative 24; Mismatches 49; Indels 13; Gaps 4;

QY 12 CVGCPRIPTNSPELEETLTHITKLNAENNATFYFKIDNVKARVQVWAGKKYFIDFVA 71

```

```

DB 154 CPDCSPIDLNSPALSAAETSLAKFNKSPSKY-ELVKTKANNQWSPAYVEVLI 212
QY 72 RETTCSK-----ESNEELTESCETKKGQSLDCNVAEYVVPWEKKIYPTVNCQ 119
DB 213 KEAPCTKQASCGLQHSDEPVGICQGSTVQSSL--RHVPLIQPKSV--TVTCE 264

RESULT 13
Q9NH95 PRELIMINARY; PRT; 148 AA.
AC Q9NH95
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Ls-cystatin.
OS Litomosoides sigmodontis.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
OC Onchocercidae; Litomosoides.
OX NCBI_TaxID=42156;
RN [1]
RP SEQUENCE FROM N.A.
RA Pfaff A.W., Hoffmann W.H., Taylor D.W., Schulz-Key H.;
RT "Characterization and immunological properties of a cysteine protease
RT inhibitor of the filarial parasite Litomosoides sigmodontis.";
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF229173; AAF35896.1; -.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00287; CYSTATIN; 1.
FT CHAIN 25 148 LS-CYSTATIN.
SQ SEQUENCE 148 AA; 16686 MW; 2950AA89CA5339C9 CRC64;

Query Match 17.2%; Score 113.5; DB 5; Length 148;
Best Local Similarity 34.1%; Pred. No. 0.0011;
Matches 30; Conservative 16; Mismatches 37; Indels 5; Gaps 3;

QY 25 ELEELTTHITKLNAENNATFYFKIDNVKARVQVWAGKKYFIDFVARETTCSENEEL 84
DB 49 EIQEMLPSTLTKVQNSNDAYHLMIKVLKVSQVWAGKKYKVCVARESDCKSSNEKI 108
QY 85 -TESCETKKGQSLD--CNVAEYVVPWE 109
DB 109 DLKTC--KKLEGHDPQIITLEVWEKWE 134

RESULT 14
Q80Y72 PRELIMINARY; PRT; 140 AA.
AC Q80Y72
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cystatin-like 1.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA TISSUE=Testicle;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Donald M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Udell T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,

```


GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:06:08 ; Search time 13.284 Seconds
(without alignments)
890.662 Million cell updates/sec

Title: US-10-661-784-1

Perfect score: 660

Sequence: 1 GKDFVQPTKICVGCPRDIP.....YVFWERKIYPTVNCQPLGM 123

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 293366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 78.*

1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	660	100.0	427	1 KGHU11	kininogen, LMW pre
2	660	100.0	644	1 KGHU11	kininogen, LMW pre
3	477	72.3	436	1 KGBOL1	kininogen, LMW I p
4	477	72.3	621	1 KGBOL1	kininogen, LMW I p
5	450	68.2	434	1 KGBOL2	kininogen, LMW II
6	450	68.2	619	1 KGBOL2	kininogen, LMW II
7	426	64.5	433	2 A28055	K-kininogen, LMW I
8	426	64.5	639	2 A28055	kininogen, LMW I p
9	409	62.0	430	2 A28055	major acute phase
10	409	62.0	430	2 A28055	T-kininogen, LMW I
11	402	60.9	423	1 KGRMT1	major acute phase
12	401	60.8	430	1 KGRMT1	T-kininogen I prec
13	136	20.6	112	1 UNBO	cystatin - bovine
14	133	20.2	91	2 S68034	T-kininogen (clone
15	133	20.2	91	2 S68035	T-kininogen (clone
16	130	19.7	127	2 S07085	cystatin C precurs
17	129	19.5	120	2 S10587	cystatin C - rat
18	128	19.4	111	2 A28793	cystatin C precurs
19	127.5	19.3	140	2 A36163	cystatin C precurs
20	127.5	19.3	146	1 UNHU	cystatin C precurs
21	125	18.9	141	2 B29632	cystatin SA precurs
22	118.5	18.0	139	1 UNCH	cystatin precursor
23	113.5	17.2	122	2 A43644	sarcocystatin A pr
24	113	17.1	141	2 J01470	cystatin S precurs
25	111.5	16.9	111	1 J02040	cystatin - chum sa
26	109.5	16.6	141	1 UNHUP2	cystatin SN precurs
27	109	16.5	141	1 UNHUP1	cystatin S precurs
28	107.5	16.3	132	2 J04918	cystatin precursor
29	107	16.2	139	2 T33740	hypothetical prote

RESULT 1

KGHUL1

kininogen, LMW precursor [validated] - human

N;Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen

N;Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen

C;Species: Homo sapiens (man)

C;Date: 06-Jul-1982 #sequence revision 27-Nov-1985 #text change 08-Dec-2000

C;Accession: A01280; B25276; A27900; A27699; A31905; A34030

R;Ohkubo, I.; Kurachi, K.; Takasawa, T.; Shiohara, H.; Sasaki, M.

Biochemistry 23, 5631-5697, 1984

A;Title: Isolation of a human cDNA for alpha-2-thiol proteinase inhibitor and its identi

A;Reference number: A90490; MUID:85122621; PMID:6441591

A;Accession: A01280

A;Molecule type: mRNA

A;Residues: 1-427 <OHK>

A;Cross-references: GB:K02566; NID:gl77889; PIDN:AAA35497.1; PID:gl77890

R;Takagaki, Y.; Kitamura, N.; Nakanishi, S.

J. Biol. Chem. 260, 8601-8609, 1985

A;Title: Cloning and sequence analysis of cDNAs for human high molecular weight and low

A;Reference number: A92544; MUID:85234582; PMID:2989293

A;Accession: B25276

A;Molecule type: mRNA

A;Residues: 1-427 <TAK>

A;Cross-references: GB:M11437; NID:gl86751; PIDN:AAB59551.1; PID:g3866853

R;Lottspeich, F.; Kellermann, J.; Henschen, A.; Rauth, G.; Mueller-Esterl, W.

in Kinins IV, part A, Greenbaum, L.M., and Margolius, H.S., eds., pp.91-95, Plenum, New

A;Title: Amino acid sequence of the light chain of human low molecular mass kininogen.

A;Reference number: A27900

A;Accession: A27900

A;Molecule type: protein

A;Residues: 390-427 <LOT>

R;Mindrou, T.; Carretero, O.A.; Proud, D.; Walz, D.; Scicli, A.G.

Biochem. Biophys. Res. Commun. 152, 519-526, 1988

A;Title: A new kinin moiety in human plasma kininogens.

A;Reference number: A27699; MUID:88209021; PMID:3365237

A;Accession: A27699

A;Molecule type: protein

A;Residues: 380-389 <MIN>

R;Maeda, H.; Matsumura, Y.; Kato, H.

J. Biol. Chem. 263, 16051-16054, 1988

A;Title: Purification and identification of [hydroxyprolyl(3)]-lysyl-bradykinin in ascitic fluid

A;Reference number: A31905; MUID:89034061; PMID:3182782

A;Accession: A31905

A;Molecule type: protein

A;Residues: 381-389 <MAE>

R;Sasaguri, M.; Ikeda, M.; Ideishi, M.; Arakawa, K.

Biochem. Biophys. Res. Commun. 150, 511-516, 1988

A;Title: Identification of [hydroxyproline(3)]-lysyl-bradykinin released from human plas

A;Reference number: A34030; MUID:88106632; PMID:3337729

A;Accession: A34030

A;Molecule type: protein

A;Residues: 380-389 <SAS>

R; Kitamura, N.; Kitagawa, H.; Fukushima, D.; Takagaki, Y.; Miyata, T.; Nakanishi, S.
 J. Biol. Chem. 260, 8610-8617, 1985
 A; Title: Structural organization of the human kininogen gene and a model for its evolution
 A; Reference number: A92545; MUID: 85234583; PMID: 2989294
 A; Contents: annotation, gene organization
 R; Pierce, J.V.
 Fed. Proc. 27, 52-57, 1968
 A; Title: Structural features of plasma kinins and kininogens.
 A; Reference number: A91455; MUID: 90255622; PMID: 4952632
 A; Contents: annotation, bradykinin
 C; Comment: The LMW kininogen precursor is produced from the same gene as the HMW form (see C; Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the C; Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, a xypoline residue is present in the kininogen prior to the release of bradykinin.
 C; Genetics:
 A; Gene: GDB:KNG
 A; Cross-references: GDB:125256; OMIM:228960
 A; Map position: 3q27-3q27
 A; Introns: 65/3, 102/3, 131/1, 188/3, 224/3, 253/1, 310/3, 346/3, 375/3, 401/3
 C; Superfamily: kininogen; cystatin homology
 C; Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; glycosylation
 F; 1-18/Domain: signal sequence #status predicted <SIG>
 F; 19-427/Product: LMW prokininogen (kininogen I) #status predicted <MAT>
 F; 19-389, 390-427/Product: LMW kininogen II #status predicted <MAT2>
 F; 19-379/Product: LMW kininogen heavy chain #status predicted <HCH>
 F; 19-131/Domain: cystatin homology <CY1>
 F; 142-253/Domain: cystatin homology <CY2>
 F; 264-375/Domain: cystatin homology <CY3>
 F; 380-389/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>
 F; 381-389/Product: bradykinin (kallidin I) #status experimental <BDY>
 F; 390-427/Product: LMW kininogen light chain #status experimental <LCH>
 F; 19/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status predicted
 F; 28-407, 83-94, 107-126, 142-145, 206-218, 229-248, 264-267, 328-340, 351-370/Disulfide bonds:
 F; 48, 469, 505, 294/Binding site: carbonylate (Asn) (covalent) #status predicted
 F; 379-380/Cleavage site: Met-Lys (kallikrein) #status experimental
 F; 383/Modified site: 4-hydroxyproline (Pro) (partial) #status experimental
 F; 389-390/Cleavage site: Arg-Ser (kallikrein) #status experimental
 F; 401/Binding site: carbonylate (Thr) (covalent) #status absent

Query Match 100.0%; Score 660; DB 1; Length 427;
 Best Local Similarity 100.0%; Pred. No. 3.3e-55;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVPPKICVCGPRDPTNSPELEETLTITTKLAENNAATFYKIDNVKKARVQV 60
 DB 253 GKDFVPPKICVCGPRDPTNSPELEETLTITTKLAENNAATFYKIDNVKKARVQV 312
 QY 61 AGKKYFIDFVARETTCKSKSNEELTESCTKLGQSLDCNAEYVVPWEKKIYPTVNCOP 120
 DB 313 AGKKYFIDFVARETTCKSKSNEELTESCTKLGQSLDCNAEYVVPWEKKIYPTVNCOP 372
 QY 121 LGM 123
 DB 373 LGM 375

RESULT 2
 KGHU1
 N; kininogen, HMW precursor [validated] - human
 N; Alternate names: alpha-2-thiol proteinase inhibitor; prokininogen; prokininogen
 N; Contains: bradykinin (kallidin I); HMW kininogen I; HMW kininogen II; low molecular weight
 C; Species: Homo sapiens (man)
 C; Date: 28-May-1986 #sequence_revision 28-May-1986 #text_change 08-Dec-2000
 C; Accession: A01279; A25276; S32422; A91153; A24871; A27899; A27699; A31905; S02
 R; Ohkubo, I.; Kurachi, K.; Takasawa, T.; Shiohara, H.; Sasaki, M.
 Biochemistry 23, 5691-5697, 1984
 A; Title: Isolation of a human cDNA for alpha-2-thiol proteinase inhibitor and its identification
 A; Reference number: A90490; MUID: 85122621; PMID: 6441591
 A; Accession: A01279
 A; Molecule type: mRNA
 A; Residues: 1-389 <OHK>
 A; Cross-references: GB:K02566; NID: g177889
 R; Takagaki, Y.; Kitamura, N.; Nakanishi, S.

J. Biol. Chem. 260, 8601-8609, 1985
 A; Title: Cloning and sequence analysis of cDNAs for human high molecular weight and low molecular weight kininogen
 A; Reference number: A92544; MUID: 85234582; PMID: 2989293
 A; Accession: A25276
 A; Molecule type: mRNA
 A; Residues: 1-592, '1', 594-644 <TAK>
 A; Cross-references: GB:M11437; NID: g186751; PIDN: AAB59550.1; PID: g386852
 R; Auerwald, E.A.; Roessler, D.; Mentele, R.; Assfalg-Machleidt, I.
 FEBS Lett. 321, 93-97, 1993
 A; Title: Cloning, expression and characterization of human kininogen domain 3.
 A; Reference number: S32422; MUID: 93223854; PMID: 8467916
 A; Accession: S32422
 A; Molecule type: mRNA
 A; Residues: 'ANSM', 253-377 <AUE>
 A; Note: differences are due to known cloning artifacts
 R; Lottspeich, F.; Kellermann, J.; Henschen, A.; Foersts, B.; Muller-Esterl, W.
 Eur. J. Biochem. 152, 307-314, 1985
 A; Title: The amino acid sequence of the light chain of human high-molecular-mass kininogen
 A; Reference number: A91153; MUID: 86030270; PMID: 4054110
 A; Accession: A91153
 A; Molecule type: protein
 A; Residues: 379-644 <LOR>
 A; Note: the bradykinin sequence preceding the light chain sequence was not determined in R; Kellermann, J.; Lottspeich, F.; Henschen, A.; Mueller-Esterl, W.
 Eur. J. Biochem. 154, 471-478, 1986
 A; Title: Completion of the primary structure of human high-molecular-mass kininogen. The
 A; Reference number: A24871; MUID: 86108361; PMID: 3484703
 A; Accession: A24871
 A; Molecule type: protein
 A; Residues: '2', 20-380 <KEL1>
 R; Kellermann, J.; Lottspeich, F.; Henschen, A.; Mueller-Esterl, W.
 in Kinins IV, Greenbaum, L.M., and Margolius, H.S., ed., pp. 85-89, Plenum Press, New York
 A; Title: Amino acid sequence of the light chain of human high molecular mass kininogen.
 A; Reference number: A27899
 A; Accession: A27899
 A; Molecule type: protein
 A; Residues: 379-389, 'K', 390-407, 'Q', 409-644 <KEL2>
 R; Miroslav, T.; Carretero, O.A.; Proud, D.; Walz, D.; Scicli, A.G.
 Biochem. Biophys. Res. Commun. 152, 519-526, 1988
 A; Title: A new kinin moiety in human plasma kininogens.
 A; Reference number: A27699; MUID: 88209021; PMID: 3365237
 A; Accession: A27699
 A; Molecule type: protein
 A; Residues: 380-389 <MIN>
 R; Maeda, H.; Matsumura, Y.; Kato, H.
 J. Biol. Chem. 263, 16051-16054, 1988
 A; Title: Purification and identification of [hydroxyprolyl(3)]bradykinin in ascitic fluid
 A; Reference number: A31905; MUID: 89034061; PMID: 3182782
 A; Accession: A31905
 A; Molecule type: protein
 R; Sasaguri, M.; Ikeda, M.; Ideishi, M.; Arakawa, K.
 Biochem. Biophys. Res. Commun. 150, 511-516, 1988
 A; Title: Identification of [hydroxyproline(3)]-lysyl-bradykinin released from human plasma
 A; Reference number: A34030; MUID: 88106632; PMID: 3337729
 A; Accession: A34030
 A; Molecule type: protein
 A; Residues: 380-389 <SAS>
 R; Lenaric, B.; Gabrijelcic, D.; Rozman, B.; Drobnic-Kosorok, M.; Turk, V.
 Biol. Chem. Hoppe-Seyler 369, 257-261, 1988
 A; Title: Human cathepsin B and cysteine proteinase inhibitors (CPis) in inflammatory and
 A; Reference number: S02482; MUID: 89076517; PMID: 3264507
 A; Accession: S02482
 A; Molecule type: protein
 A; Residues: 1-19; 189-192; 310-314; 381-389 <LEN1>
 R; Kato, H.; Matsumura, Y.; Maeda, H.
 FEBS Lett. 232, 252-254, 1988
 A; Title: Isolation and identification of hydroxyproline analogues of bradykinin in human
 A; Reference number: A61495; MUID: 88211869; PMID: 3366244
 A; Accession: A61495
 A; Molecule type: protein
 A; Residues: 380-389 <KAT1>
 A; Experimental source: urine

A>Note: this peptide had Pro-383 modified to 4-hydroxyproline

A:Accession: B61495
 A:Molecule type: protein
 A:Residues: 381-389 <KAT2>
 A:Experimental source: urine
 A>Note: this peptide had Pro-383 modified to 4-hydroxyproline
 A:Accession: C61495
 A:Molecule type: protein
 A:Residues: 380-389 <KAT3>
 R:Lenarcic, B.; Krasovec, M.; Ritonja, A.; Olafsson, I.; Turk, V.
 FEBS Lett. 280, 211-215, 1991
 A>Title: Inactivation of human cystatin C and kininogen by human cathepsin D.
 A:Reference number: S14303; MUID:91192133; PMID:2013314
 A:Accession: S14447
 A:Molecule type: protein
 A:Residues: 264-359, 'N', 361-375 <LEN2>
 R:Littell, S.S.; Johnson, D.A.
 Biochem. J. 307, 341-346, 1995
 A>Title: Human mast cell tryptase isoforms: separation and examination of substrate-specificity
 A:Reference number: S55239; MUID:95251593; PMID:7733867
 A:Accession: S55239
 A:Molecule type: protein
 A:Residues: 450-452, 'X', 454, 'X', 456 <JIT>
 R:Straczek, J.; Maachi, F.; le Nguyen, D.; Becchi, M.; Heulin, M.H.; Nabat, P.; Bellevil
 FEBS Lett. 373, 207-211, 1995
 A>Title: Purification from human plasma of a tetrapeptide that potentiates insulin-like
 A:Reference number: S68059; MUID:96033974; PMID:7589467
 A:Accession: S68059
 A:Molecule type: protein
 A:Residues: 431-434 <STR>
 R:Kitamura, N.; Kitagawa, H.; Fukushima, D.; Takagaki, Y.; Miyata, T.; Nakanishi, S.
 J. Biol. Chem. 260, 8610-8617, 1985
 A>Title: Structural organization of the human kininogen gene and a model for its evolution
 A:Reference number: A92545; MUID:85234583; PMID:2989294
 A:Contents: annotation; gene organization
 R:Pierce, J.V.
 Fed. Proc. 27, 52-57, 1968
 A>Title: Structural features of plasma kinins and kininogens.
 A:Reference number: A91455; MUID:90255622; PMID:4952632
 A:Contents: annotation; bradykinin
 C:Comment: The HMW kininogen precursor and the LMW form are produced from the same gene
 C:Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
 C:Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is impo
 C:Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i
 xyproline residue is present in the kininogen prior to the release of bradykinin.
 C:Genetics:
 A:Gene: GDB:KNG
 A:Cross-references: GDB:125256; OMIM:228960
 A:Map position: 3q27-3q27
 A:Introns: 65/3; 102/3; 131/1; 188/3; 224/3; 253/1; 310/3; 346/3; 375/3
 A:Superfamily: kininogen; cystatin homology
 C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; dupl
 F:1-18/Domain: signal sequence #status experimental <SIG>
 F:19-644/Product: HMW kininogen I (prokininogen) #status experimental <MAT1>
 F:19-379,390-644/Product: HMW kininogen II #status experimental <MAT2>
 F:19-379/Domain: HMW kininogen heavy chain #status experimental <HCH>
 F:19-131/Domain: cystatin homology <CY1>
 F:142-253/Domain: cystatin homology <CY2>
 F:264-375/Domain: cystatin homology <CY3>
 F:380-389/Product: lysyl-bradykinin (kallidin II) #status experimental <XBDY>
 F:381-389/Product: bradykinin (kallidin I) #status experimental <BDY>
 F:390-644/Domain: HMW kininogen light chain #status experimental <LCH>
 F:421-510/Region: glycine/histidine/lysine-rich 30-residue repeats
 F:19/Modified site: low molecular weight growth promoting factor #status experimental <GF
 F:28-614,83-94,107-126,145-206-218,229-248,264-267,328-340,351-370/Disulfide bonds:
 F:48/Binding site: carboxylate (Asn) (covalent) #status absent
 F:169,205,294/Binding site: carboxylate (Asn) (covalent) #status experimental
 F:379-380/Cleavage site: Met-Lys (kallikrein) #status experimental
 F:383/Modified site: 4-hydroxyproline (Pro) (partial) #status experimental
 F:389-390/Cleavage site: Arg-Ser (kallikrein) #status experimental
 F:401,533,542,546,557,571,593,628/Binding site: carboxylate (Thr) (covalent) #status ex
 F:577/Binding site: carboxylate (Ser) (covalent) #status experimental

Query Match 100.0%; Score 660; DB 1; Length 644;
 Best Local Similarity 100.0%; Pred. NO. 5.1e-55;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKKARQVQV 60
 |||||
 DB 253 GKDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKKARQVQV 312
 |||||

QY 61 AGKKYFIDFVARETTCESNEELTESCTKKLGSLDCNAEYVYVVPWEKKIYPTVNCQP 120
 |||||
 DB 313 AGKKYFIDFVARETTCESNEELTESCTKKLGSLDCNAEYVYVVPWEKKIYPTVNCQP 372
 |||||

QY 121 LGM 123
 ||||
 DB 373 LGM 375

RESULT 3

KGBOLI

kininogen, LMW I precursor - bovine

N/Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen

N/Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen

C/Species: Bos primigenius taurus (cattle)

C/Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 22-Jun-1999

C/Accession: A01283

R:Nawa, H.; Kitamura, N.; Hirose, T.; Asai, M.; Inayama, S.; Nakanishi, S.

Proc. Natl. Acad. Sci. U.S.A. 80, 90-94, 1983

A>Title: Primary structures of bovine liver low molecular weight kininogen precursors an

A/Reference number: A93984; MUID:83117859; PMID:6572010

A/Accession: A01283

A:Molecule type: mRNA

A:Residues: 1-436 <NAW>

A:Cross-references: GB:J00010; GB:V00426; NID:G163256; PID:AAA30604.1; PID:G163257

C/Comment: The LMW kininogen precursor is produced from the same gene as the HMW form as

C/Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the

C/Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i

xyproline residue is present in the kininogen prior to the release of bradykinin.

C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; glyc

F:1-18/Domain: signal sequence #status predicted <SIG>

F:19-436/Product: LMW kininogen I #status predicted <MAT>

F:19-378/Product: LMW kininogen I heavy chain #status predicted <HCH>

F:19-130/Domain: cystatin homology <CY1>

F:141-252/Domain: cystatin homology <CY2>

F:263-374/Domain: cystatin homology <CY3>

F:379-388/Product: lysyl-bradykinin (kallidin II) #status predicted <XBDY>

F:380-388/Product: bradykinin (kallidin I) #status predicted <BDY>

F:389-436/Product: LMW kininogen I light chain #status experimental <LCH>

F:19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status predicted

F:27-406,82-93,106-125,141-144,205-217,228-247,263-286,327-339,350-369/Disulfide bonds:

F:47,87,168,169,197,204/Binding site: carboxylate (Asn) (covalent) #status predicted

F:378-379/Cleavage site: Met-Lys (kallikrein) #status predicted

F:382/Modified site: 4-hydroxyproline (Pro) #status predicted

F:388-389/Cleavage site: Arg-Ser (kallikrein) #status predicted

Query Match 72.3%; Score 477; DB 1; Length 436;

Best Local Similarity 71.9%; Pred. NO. 9.2e-38;

Matches 87; Conservative 14; Mismatches 20; Indels 0; Gaps 0;

QY 2 KDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKKARQVQV 61
 |||||
 DB 253 KDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKKARQVQV 312
 |||||

QY 62 GKXYFIDFVARETTCESNEELTESCTKKLGSLDCNAEYVYVVPWEKKIYPTVNCQP 121
 |||||
 DB 313 GLKYSIVFARETTCESNEELTESCTKKLGSLDCNAEYVYVVPWEKKIYPTVNCQP 372
 |||||

QY 122 G 122
 ||||
 DB 373 G 373

Db 370 LG 371

RESULT 6

K60H2

kininogen, HMW II precursor - bovine
N:Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen
N:Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C:Species: Bos primigenius taurus (cattle)
C>Date: 14-Nov-1983 #sequence revision 14-Nov-1983 #text_change 22-Jun-1999
A:Accession: A01282; A91923; A91941; A91938; B29559
R:Kitamura, N.; Takagaki, Y.; Furuto, S.; Tanaka, T.; Nawa, H.; Nakanishi, S.
Nature 305, 545-549, 1983
A>Title: A single gene for bovine high molecular weight and low molecular weight kininogen
A:Reference number: A93317; PMID:84014106; PMID:6571699
A:Accession: A01282
A:Molecule type: mRNA
A:Residues: 1-619 <KT>
A:Cross-references: GB:V01492; GB:K01758; NID:6493; PIDN:CAA24736.1; PID:9494
R:Kato, H.; Nagasawa, S.; Suzuki, T.
J. Biochem. 67, 313-323, 1970
A>Title: Studies on the structure of bovine kininogen: cleavages of disulfide bonds and
A:Reference number: A91923; PMID:70180420; PMID:4986212
A:Accession: A91923
A:Molecule type: protein
A:Residues: 378-391 <KAT>
R:Han, Y.N.; Kato, H.; Iwanaga, S.; Suzuki, T.
J. Biochem. 79, 1201-1222, 1976
A>Title: Primary structure of bovine plasma high-molecular-weight kininogen. The amino a
A:Reference number: A91941; PMID:76260155; PMID:956151
A:Accession: A91941
A:Molecule type: protein
A:Residues: 387-455 <HAN>
A>Note: 398-Pro, 401-Val, and 455-Lys were also found
R:Han, Y.N.; Komiya, M.; Iwanaga, S.; Suzuki, T.
J. Biochem. 77, 55-68, 1975
A>Title: Studies on the primary structure of bovine high-molecular-weight kininogen. Ami
A:Reference number: A91938; PMID:75170265; PMID:1169237
A:Accession: A91938
A:Molecule type: protein
A:Residues: 456-496 <HA2>
R:Sueyoshi, T.; Miyata, T.; Hashimoto, N.; Kato, H.; Hayashida, H.; Miyata, T.; Iwanaga,
J. Biol. Chem. 262, 2768-2779, 1987
A>Title: Bovine high molecular weight kininogen. The amino acid sequence, positions of c
A:Reference number: A92627; PMID:87137530; PMID:3546295
A:Accession: B29559
A:Molecule type: protein
A:Residues: 'Z', 20-104, 'E', 106-256, 'XX', 257-376 <SUB>
R:Lottspeich, F.; Kellermann, J.; Henschen, A.; Foerisch, B.; Muller-Esterl, W.
Eur. J. Biochem. 152, 307-314, 1985
A>Title: The amino acid sequence of the light chain of human high-molecular-mass kininog
A:Reference number: A91153; PMID:86030270; PMID:4054110
A:Contents: annotation; bovine cleavage sites; bovine carbohydrate binding sites
R:Sueyoshi, T.; Miyata, T.; Kato, H.; Iwanaga, S.
Seikagaku 56, 808, 1984
A>Title: Disulfide bonds in bovine HMW kininogens.
A:Reference number: A94300
A:Contents: annotation; disulfide bonds
A>Note: article in Japanese
C:Comment: the HMW kininogen precursor is produced from the same gene as the LMW form as
C:Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
C:Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is impo
C:Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, a
xyproline residue is present in the kininogen prior to the release of bradykinin.
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; dupl
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-619/Product: HMW kininogen II #status predicted <NAT>
F:19-376/Product: HMW kininogen II heavy chain #status experimental <HCH>
F:19-130/Domain: cystatin homology <CY1>
F:141-252/Domain: cystatin homology <CY2>
F:261-372/Domain: cystatin homology <CY3>
F:377-386/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>

F:378-386/Product: bradykinin (kallidin I) #status experimental <BDY>
F:387-619/Product: HMW kininogen II light chain #status experimental <LCH>
F:418-488/Region: glycine/histidine/lysine-rich
F:19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status experimen
F:27-589 82-93 106-125 141-144 205-217 228-247 261-264 325-337 348-367/Disulfide bonds:
F:47/Binding site: carbohydrate (Asn) (covalent) #status absent
F:87,168,169,204,280/Binding site: carbohydrate (Asn) (covalent) #status experimental
F:136/Binding site: carbohydrate (Thr) (covalent) (partial) #status experimental
F:197/Binding site: carbohydrate (Asn) (covalent) (partial) #status experimental
F:376-377/Cleavage site: Met-Lys (kallikrein) #status experimental
F:380/Modified site: 4-hydroxyproline (Pro) #status predicted
F:386-387/Cleavage site: Arg-Ser (kallikrein) #status experimental
F:396,400,404,510/Binding site: carbohydrate (Ser) (covalent) #status experimental
F:397,398,518,522,534,546,551,569/Binding site: carbohydrate (Thr) (covalent) #status ex
F:496-497/Cleavage site: Arg-Thr (kallikrein) #status experimental

Query Match 68.2%; Score 450; DB 1; Length 619;
Best Local Similarity 68.9%; Pred. No. 5.1e-35;
Matches 84; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

QY 1 GKDFVOPPTKICVGCPRDPTNSPELEETLTTHITKLNANNATFYFKIDNVKKARQVY 60
DB 252 GEDFL--PMWCVGCCPKIPVDSPLDEALNHSIAKLNAEHDGTFFYKIDTVKKATVQVY 309

QY 61 AGKVFIDFVARETTCKSKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTVNCOP 120
DB 310 GGLKYSIVFIARETTCKSKSNEELTKSCHEINHGQLLHCDANVYVVPWEKKYPTVNCOP 369

QY 121 LG 122
DB 370 LG 371

RESULT 7
K-kininogen, LMW I precursor - rat
A:Accession: A28055
C:Species: Rattus norvegicus (Norway rat)
C>Date: 20-Jun-1989 #sequence_revision 20-Jun-1989 #text_change 15-Nov-1996
A>Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and
R:Furuto-Kato, S.; Matsumoto, A.; Kitamura, N.; Nakanishi, S.
J. Biol. Chem. 260, 12054-12059, 1985
A>Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and
A:Reference number: A92496; PMID:86008264; PMID:2413018
A:Accession: A28055
A:Molecule type: mRNA
A:Residues: 1-433 <FUR>
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-433/Product: K-kininogen, LMW I #status predicted <NAT>
F:19-131/Domain: cystatin homology <CY1>
F:142-253/Domain: cystatin homology <CY2>
F:264-375/Domain: cystatin homology <CY3>

Query Match 64.5%; Score 426; DB 2; Length 433;
Best Local Similarity 65.0%; Pred. No. 6.6e-33;
Matches 80; Conservative 14; Mismatches 29; Indels 0; Gaps 0;

QY 1 GKDFVOPPTKICVGCPRDPTNSPELEETLTTHITKLNANNATFYFKIDNVKKARQVY 60
DB 253 GDDLPELLPDCPCGPRNIPVDSPELKEALGHSIAQLNANNHTFFYKIDTVKKATSOVY 312

QY 61 AGKVFIDFVARETTCKSKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTVNCOP 120
DB 313 AGTKVIEFIARETTCKSKSNEELTADCTETKLGQSLNCANVYVMPWENKVVPTVKCY 372

QY 121 LGM 123
DB 373 LDM 375

RESULT 8

A25486
kininogen, HMW I precursor - rat
N:Contains: bradykinin
C:Species: Rattus norvegicus (Norway rat)
C>Date: 08-Mar-1989 #sequence_revision 08-Mar-1989 #text_change 15-Nov-1996
C/Accession: A25486
R:Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 262, 2190-2198, 1987
A>Title: Differing expression patterns and evolution of the rat kininogen gene family.
A:Reference number: A92625; MUID:87137443; PMID:3029068
A:Accession: A25486
A:Molecule type: mRNA
A:Residues: 1-639 <KIT>
A>Note: the authors translated the codon CAA for residue 347 as Asn
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-639/Product: kininogen, HMW I #status predicted <MAT>
F:19-131/Domain: cystatin homology <CY1>
F:142-253/Domain: cystatin homology <CY2>
F:264-375/Domain: cystatin homology <CY3>
Query Match 64.5%; Score 426; DB 2; Length 639;
Best Local Similarity 65.0%; Pred. No. 1e-32;
Matches 80; Conservative 14; Mismatches 29; Indels 0; Gaps 0;
QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLAENNATFYKIDNVKKARVQV 60
DB 253 GDDLFSLLPKKCGCPKNIPVDSPELKEALGHSIAQLNAQNHLYFKIDTVKKATSQV 312
QY 61 AGKKYFIDFVARETTCSKESNEELTESCTKLGOSLDCNAEVVYVWPKKIYPTVNCOP 120
DB 313 AGTKYVIEFIARETNCSTQNTLTADCTKHLGQSLNCNANVYRPWENKVVPTVRCQA 372
QY 121 LGM 123
DB 373 LDM 374
RESULT 9
A23897
Major acute phase alpha-1 protein (version 2) - rat
C:Species: Rattus norvegicus (Norway rat)
C>Date: 19-May-1989 #sequence_revision 19-May-1989 #text_change 20-Aug-1999
C/Accession: A23897; B23897
R:Anderson, K.P.; Heath, B.C.
J. Biol. Chem. 260, 12065-12071, 1985
A>Title: The relationship between rat major acute phase protein and the kininogens.
A:Reference number: A23897; MUID:86008266; PMID:2413019
A:Accession: A23897
A:Molecule type: protein
A:Residues: 1-14 <AND1>
A:Accession: B23897
A:Molecule type: mRNA
A:Residues: 5-430 <AND2>
A/Cross-references: GB:M11661; NID:G205307; PID:AAA41570.1; PID:G205308
A>Note: the authors translated the codon CTC for residue 410 as Arg, CTA for residue 415
C:Superfamily: kininogen; cystatin homology
F:19-130/Domain: cystatin homology <CY1>
F:141-252/Domain: cystatin homology <CY2>
F:263-374/Domain: cystatin homology <CY3>
Query Match 62.0%; Score 409; DB 2; Length 430;
Best Local Similarity 61.8%; Pred. No. 2.7e-31;
Matches 76; Conservative 15; Mismatches 33; Indels 0; Gaps 0;
QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLAENNATFYKIDNVKKARVQV 60
DB 252 GDDLFSLLPKKCGCPKNIPVDSPELKEALGHSIAQLNAQNHLYFKIDTVKKATSQV 311
QY 61 AGKKYFIDFVARETTCSKESNEELTESCTKLGOSLDCNAEVVYVWPKKIYPTVNCOP 120
DB 312 AGTKYVIEFIARETNCSTQNTLTADCTKHLGQSLNCNANVYRPWENKVVPTVRCQA 371
Query Match 62.0%; Score 409; DB 2; Length 430;
Best Local Similarity 61.8%; Pred. No. 2.7e-31;
Matches 76; Conservative 15; Mismatches 33; Indels 0; Gaps 0;
QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLAENNATFYKIDNVKKARVQV 60
DB 252 GDDLFSLLPKKCGCPKNIPVDSPELKEALGHSIAQLNAQNHLYFKIDTVKKATSQV 311
QY 61 AGKKYFIDFVARETTCSKESNEELTESCTKLGOSLDCNAEVVYVWPKKIYPTVNCOP 120
DB 312 AGTKYVIEFIARETNCSTQNTLTADCTKHLGQSLNCNANVYRPWENKVVPTVRCQA 371

QY 121 LGM 123
DB 372 LDM 374
RESULT 10
B28055
T-kininogen, LMW II precursor - rat
C:Species: Rattus norvegicus (Norway rat)
C>Date: 20-Jun-1989 #sequence_revision 20-Jun-1989 #text_change 12-Dec-1997
C/Accession: B28055; E25486; E28526; C28526
R:Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 262, 2190-2198, 1987
A>Title: Differing expression patterns and evolution of the rat kininogen gene family.
A:Reference number: A92625; MUID:87137443; PMID:3029068
A:Accession: E25486
A:Molecule type: DNA
A:Residues: 375-430 <KIT>
R:Enjyoji, K.; Kato, H.; Hayaashi, I.; Oh-ishi, S.; Iwanaga, S.
J. Biol. Chem. 263, 973-979, 1988
A>Title: Purification and characterization of rat T-kininogens isolated from plasma of a
A:Reference number: A92729; MUID:88087226; PMID:3121623
A:Accession: B28526
A:Molecule type: protein
A:Residues: 'B', 20-25, 'MD', 28-48, 376-430 <ENJ>
A:Accession: C28526
A:Molecule type: protein
A:Residues: 'B', 20-48, 376-388, 'R', 390-419, 'ER', 423-430 <EN2>
C:Superfamily: kininogen; cystatin homology
C:Keywords: glycoprotein; pyroglutamic acid
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-430/Product: T-kininogen, LMW II #status experimental <MAT>
F:19-130/Domain: cystatin homology <CY1>
F:141-252/Domain: cystatin homology <CY2>
F:263-374/Domain: cystatin homology <CY3>
F:19/Modified site: pyroglutamic acid (Gln) (in mature form) #status experiment
F:82,126,168,204,326/Binding site: carboxylate (Asn) (covalent) #status predicted
F:83-94,107-125,141-144,205-217,228-247,263-286,327-339,350-369/Disulfide bonds: #status
Query Match 62.0%; Score 409; DB 2; Length 430;
Best Local Similarity 61.8%; Pred. No. 2.7e-31;
Matches 76; Conservative 15; Mismatches 32; Indels 0; Gaps 0;
QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLAENNATFYKIDNVKKARVQV 60
DB 252 GDDLFSLLPKKCGCPKNIPVDSPELKEALGHSIAQLNAQNHLYFKIDTVKKATSQV 311
QY 61 AGKKYFIDFVARETTCSKESNEELTESCTKLGOSLDCNAEVVYVWPKKIYPTVNCOP 120
DB 312 AGTKYVIEFIARETNCSTQNTLTADCTKHLGQSLNCNANVYRPWENKVVPTVRCQA 371
QY 121 LGM 123
DB 372 LDM 374
RESULT 11
XGRFM
Major acute phase alpha-1 protein precursor - rat (fragment)
N:Contains: bradykinin
C:Species: Rattus norvegicus (Norway rat)
C>Date: 27-Nov-1985 #sequence_revision 27-Nov-1985 #text_change 12-Apr-1996
C/Accession: A01285
R:Coile, T.; Inglis, A.S.; Roxburgh, C.M.; Howlett, G.J.; Schreiber, G.

FEBS Lett. 182, 57-61, 1985
A>Title: Major acute phase alpha-1 protein of the rat is homologous to bovine kininogen A
A:Reference number: A01285; MUID:85127561; PMID:2578992
A:Accession: A01285
A:Molecule type: mRNA
A:Residues: 1-423 <COL>
C:Comment: This plasma glycoprotein inhibits cysteine proteinases. During acute inflammation
X:
C:Superfamily: kininogen; cystatin homology
C:Keywords: bradykinin; cysteine proteinase inhibitor; duplication; glycoprotein; inflammation
F:1-11/Domain: signal sequence (fragment) #status predicted <SIG>
F:12-423/Product: major acute phase alpha-1 protein #status predicted <MAP>
F:12-123/Domain: cystatin homology <CY1>
F:134-245/Domain: cystatin homology <CY2>
F:256-357/Domain: cystatin homology <CY3>
F:371-379/Product: bradykinin #status predicted <BDY>
F:12/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status predicted
F:161,197/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 60.9%; Score 402; DB 1; Length 423;
Best Local Similarity 61.8%; Pred. No. 1.2e-30;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;

Qy 1 GKDFVQPTKICVGCPRDIPNSPELEETLTHITTKNAENNAFYFKIDNVKARVQV 60
Db 245 GDDLPELLKNCRCGPRIEIPVDSPELKEALGHSIARLNQHNHIFYFKIDTVKATSQV 304

Qy 61 AGKYFIDFVARETTCSENEELTESCTKKLGSLDCNAEVVVPWEKKIVPTVNCOP 120
Db 305 AGVIVIEFIARETNCQSKTELTADCTCKHLGSLNCNANVYMRPWENKVPTVRCOA 364

Qy 121 LGM 123
Db 365 LDM 367

RESULT 12
KGRTTI
N:kininogen I precursor - rat
N:Alternate names: 73K protein; LMW kininogen T-I
N:Contains: bradykinin; T-kinin
C:Species: Rattus norvegicus (Norway rat)
C>Date: 17-Mar-1987 #sequence revision 17-Mar-1987 #text change 22-Jun-1999
C:Accession: A01286; D25486; E0193; JQ0027; B25488; A28525; S68036
R:Furuto-Kato, S.; Matsumoto, A.; Kitamura, N.; Nakanishi, S.
J. Biol. Chem. 260, 12054-12059, 1985
A>Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and inhibitor.
A:Reference number: A92496; MUID:86008264; PMID:2413018
A:Accession: A01286
A:Molecule type: mRNA
A:Residues: 1-430 <FUR>
A:Cross-references: GB:M11883; NID:g205084; PIDN:AAA41489.1; PID:g205085
R:Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 262, 2190-2198, 1987
A>Title: Differing expression patterns and evolution of the rat kininogen gene family.
A:Reference number: A92625; MUID:87137443; PMID:3023068
A:Accession: D25486
A:Molecule type: DNA
A:Residues: 375-430 <KIT>
R:Enjiyoji, K.; Kato, H.; Hayashi, I.; Oh-ishi, S.; Iwanaga, S.
J. Biol. Chem. 263, 973-979, 1988
A>Title: Purification and characterization of rat T-kininogens isolated from plasma of a
A:Reference number: A92729; MUID:88087226; PMID:3121623
A:Accession: A28526
A:Molecule type: protein
A:Residues: 'E', 20-48; 376-430 <BNJ>
R:Kanda, S.; Sugiyama, K.; Takahashi, M.; Shumiya, S.; Tomino, S.; Nagase, S.
Jpn. J. Cancer Res. 81, 63-68, 1990
A>Title: Identification of a protein increasing in serum of Nagase analbuminemic rats be
A:Reference number: PLO193; MUID:90216390; PMID:2108948
A:Accession: PLO193
A:Molecule type: mRNA

A:Residues: 330-420, 'R', 422-429, 'P' <KAN>
R:Anderson, K.P.; Croyle, M.L.; Lingrel, J.B.
Gene 81, 119-128, 1989
A>Title: Primary structure of a gene encoding rat T-kininogen.
A:Reference number: JQ0027; MUID:90034172; PMID:2806908
A:Accession: JQ0027
A:Molecule type: DNA
A:Residues: 1-60, 'E', 62-113, 'R', 115-165, 'F', 167-178, 'TKI', 182-211, 'F', 213-256, 'S', 259-388
A:Experimental source: strain Sprague-Dawley
R:Kagayama, R.; Kitamura, N.; Okkubo, H.; Nakanishi, S.
J. Biol. Chem. 262, 2345-2355, 1987
A>Title: Differing utilization of homologous transcription initiation sites of rat K and
A:Reference number: A25488; MUID:87137465; PMID:3818598
A:Accession: B25488
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-48 <XAG>
A:Cross-references: GB:M14356; NID:g205090; PIDN:AAA41492.1; PID:g205091
R:Enjiyoji, K.; Kato, H.; Hayashi, I.; Oh-ishi, S.; Iwanaga, S.
J. Biol. Chem. 263, 965-972, 1988
A>Title: Purification and characterization of two kinds of low molecular weight kininogen
A:Reference number: A28525; MUID:88087225; PMID:3335530
A:Accession: A28525
A:Molecule type: protein
A:Residues: 376-430 <EN2>
R:Sierra, F.; Walter, R.; Vautravers, P.; Guigoz, Y.
Arch. Biochem. Biophys. 322, 333-338, 1995
A>Title: Identification of several isoforms of T-kininogen expressed in the liver of agi
A:Reference number: S68034; MUID:96032652; PMID:7574705
A:Accession: S68036
A:Molecule type: mRNA
A:Residues: 340-430 <SIE>
A:Experimental source: clone pSG17
C:Comment: At least three types of LMW kininogen precursors are present in rat plasma, t
ceding bradykinin.
C:Comment: T-kininogens contain T-kinin (I-S-bradykinin), a novel kinin isolated after t
d of an Arg or Lys, it is probably not released from its precursor by either tissue or p
C:Comment: The T-kininogens are produced in response to an inflammatory stimulant.
C:Genetics:
A:Introns: 65/3; 102/3; 130/1; 187/3; 223/2; 252/1; 309/3; 345/3; 374/3; 398/3
C:Superfamily: kininogen; cystatin homology
C:Keywords: acute phase; bradykinin; cysteine proteinase inhibitor; duplication; glycop
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-430/Product: T-kininogen I #status experimental <MAT>
F:19-130/Domain: cystatin homology <CY1>
F:141-252/Domain: cystatin homology <CY2>
F:263-374/Domain: bradykinin #status predicted <BDY>
F:378-386/Product: pyrrolidone carboxylic acid (Gln) (in mature form) #status experimen
F:19/Modified site: pyrrolidone carboxylic acid (Asn) (covalent) #status predicted
F:82, 126, 168, 204, 325/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:83-94, 107-125, 141-144, 205-217, 228-247, 263-266, 327-339, 350-369/Disulfide bonds: #status

Query Match 60.8%; Score 401; DB 1; Length 430;
Best Local Similarity 61.8%; Pred. No. 1.6e-30;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;

Qy 1 GKDFVQPTKICVGCPRDIPNSPELEETLTHITTKNAENNAFYFKIDNVKARVQV 60

Db 252 GDDLPELLKNCRCGPRIEIPVDSPELKEALGHSIARLNQHNHIFYFKIDTVKATSQV 311

Qy 61 AGKYFIDFVARETTCSENEELTESCTKKLGSLDCNAEVVVPWEKKIVPTVNCOP 120

Db 312 AGVIVIEFIARETNCQSKTELTADCTCKHLGSLNCNANVYMRPWENKVPTVRCOA 371

Qy 121 LGM 123

Db 372 LDM 374

RESULT 13

UDBO

Cystatin - bovine
N:Alternate names: thiol proteinase inhibitor

C:Species: Bos primigenius taurus (cattle)
C:Date: 28-Feb-1986 #sequence_revision 28-Feb-1986 #text_change 06-Dec-1996
C:Accession: A01271
R:Hiirado, M.; Tsunasawa, S.; Sakiyama, P.; Niinobe, M.; Fujii, S.
FEBS Lett. 186, 41-45, 1985
A:Title: Complete amino acid sequence of bovine colostrum low-M-r cysteine proteinase in
A:Reference number: A01271; MUID:85231205; PMID:3891407
A:Accession: A01271

A:Molecule type: protein
A:Residues: 1-112 <HR>
C:Superfamily: cystatin; cystatin homology
C:Keywords: colostrum; cysteine proteinase inhibitor
F:2-112/Domain: cystatin homology <CYS>
F:48-52/Region: inhibitory #status predicted
F:66-76,90-110/Disulfide bonds: #status predicted

Query Match 20.6%; Score 136; DB 1; Length 112;
Best Local Similarity 30.6%; Pred. No. 6.4e-06;
Matches 34; Conservative 22; Mismatches 31; Indels 24; Gaps 5;

QY 22 NSPELEETITHTIKLNAENNAFFPKIDNVKARVOVAGKYYFIDFVARETTCSKSN 81

DB 12 NEEGQVQALSFVAFSEFNKSGNDAYOSRVVVRARQVWGMNFDVDELGRITCTK--S 69

QY 82 BELTESC-----ETKLGQSLDCNAEVVVPWEKKIYPTVN-----CQ 119

DB 70 QANLDSCPHPNPHLKREKL-----CSFQVYVVPWMN-----TINLVKFSQ 111

RESULT 14

S68034

T-kininogen (clone pSG22) - rat (fragment)
C:Species: Rattus norvegicus (Norway rat)
C:Date: 12-Feb-1998 #sequence_revision 20-Feb-1998 #text_change 07-May-1999
C:Accession: S68034
R:Sierra, F.; Walter, R.; Vautravers, P.; Guigoz, Y.
Arch. Biochem. Biophys. 322, 333-338, 1995
A:Title: Identification of several isoforms of T-kininogen expressed in the liver of ag
A:Reference number: S68034; MUID:96032652; PMID:7574705
A:Accession: S68034

A:Molecule type: mRNA

A:Residues: 1-91 <SIE>

A:Experimental source: liver

C:Superfamily: kininogen; cystatin homology

C:Keywords: alternative splicing; cysteine proteinase inhibitor; plasma

Query Match 20.2%; Score 133; DB 2; Length 91;
Best Local Similarity 68.6%; Pred. No. 9.8e-06;
Matches 24; Conservative 3; Mismatches 9; Indels 0; Gaps 0;

QY 89 ETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 123

DB 1 ETKHGLQSLNCNANVVRPWNKVVPTVRCQALDM 35

RESULT 15

S68035

T-kininogen (clone pSG17) - rat (fragment)
C:Species: Rattus norvegicus (Norway rat)
C:Date: 12-Feb-1998 #sequence_revision 20-Feb-1998 #text_change 07-May-1999
C:Accession: S68035
R:Sierra, F.; Walter, R.; Vautravers, P.; Guigoz, Y.
Arch. Biochem. Biophys. 322, 333-338, 1995
A:Title: Identification of several isoforms of T-kininogen expressed in the liver of ag
A:Reference number: S68034; MUID:96032652; PMID:7574705
A:Accession: S68035

A:Molecule type: mRNA

A:Residues: 1-91 <SIE>

A:Experimental source: liver

C:Superfamily: kininogen; cystatin homology

C:Keywords: alternative splicing; cysteine proteinase inhibitor; plasma

Query Match 20.2%; Score 133; DB 2; Length 91;

Best Local Similarity 68.6%; Pred. No. 9.8e-06;
Matches 24; Conservative 3; Mismatches 8; Indels 0; Gaps 0;

QY 89 ETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 123

DB 1 ETKHGLQSLNCNANVVRPWNKVVPTVRCQALDM 35

Search completed: September 24, 2004, 14:10:48
Job time: 14.284 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:08:41 : Search time 43.296 Seconds
(without alignments)
913.519 Million cell updates/sec

Title: US-10-661-784-1
Perfect score: 660
Sequence: 1 GKDFVQPTKICVGRDIP.....YVWPWEKKIYPTVNCQPLGM 123

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1349238 seqs, 321558718 residues

Total number of hits satisfying chosen parameters: 1349238

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications AA:*
1: /cgn2_6/ptodata/1/pubpaa/US07_PUBCOMB.pep.*
2: /cgn2_6/ptodata/1/pubpaa/PCT_NEW_PUB.pep.*
3: /cgn2_6/ptodata/1/pubpaa/US06_NEW_PUB.pep.*
4: /cgn2_6/ptodata/1/pubpaa/US06_PUBCOMB.pep.*
5: /cgn2_6/ptodata/1/pubpaa/US07_NEW_PUB.pep.*
6: /cgn2_6/ptodata/1/pubpaa/PCTUS_PUBCOMB.pep.*
7: /cgn2_6/ptodata/1/pubpaa/US08_NEW_PUB.pep.*
8: /cgn2_6/ptodata/1/pubpaa/US08_PUBCOMB.pep.*
9: /cgn2_6/ptodata/1/pubpaa/US09_PUBCOMB.pep.*
10: /cgn2_6/ptodata/1/pubpaa/US09B_PUBCOMB.pep.*
11: /cgn2_6/ptodata/1/pubpaa/US09C_PUBCOMB.pep.*
12: /cgn2_6/ptodata/1/pubpaa/US09_NEW_PUB.pep.*
13: /cgn2_6/ptodata/1/pubpaa/US10A_PUBCOMB.pep.*
14: /cgn2_6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	560	100.0	390	15	US-10-162-335-82
2	560	100.0	398	15	US-10-162-335-70
3	560	100.0	427	10	US-09-919-039-29
4	560	100.0	615	15	US-10-162-335-72
5	560	100.0	644	15	US-10-162-335-74
6	560	100.0	644	15	US-10-162-335-84
7	402	60.9	424	14	US-10-316-253-217
8	401	60.8	430	14	US-10-316-253-215
9	169	25.6	178	9	US-09-969-834-1
10	165.5	25.1	145	14	US-10-329-428-2
11	165.5	25.1	167	10	US-09-746-783-197
12	138.5	21.0	121	9	US-09-775-932-14
13	138.5	21.0	128	9	US-09-775-932-12
14	138.5	21.0	149	9	US-09-940-497-2
15	136	20.6	112	8	US-08-849-303-16

16	136	20.6	112	16	US-10-655-136-16
17	135	20.5	118	9	US-09-775-932-24
18	130	19.7	127	16	US-08-849-303-19
19	130	19.7	127	16	US-10-655-136-19
20	129.5	19.6	140	14	US-10-376-564-46
21	129.5	19.6	140	14	US-10-376-564-48
22	129.5	19.6	317	12	US-10-210-172-82
23	129.5	19.6	345	12	US-10-210-172-86
24	129.5	19.6	356	12	US-10-210-172-84
25	129.5	19.6	369	12	US-10-210-172-78
26	129.5	19.6	369	12	US-10-210-172-80
27	129.5	19.6	382	12	US-10-315-664-93
28	129.5	19.6	382	12	US-09-978-360A-425
29	128	19.4	111	8	US-08-849-303-26
30	128	19.4	111	16	US-10-655-136-26
31	127.5	19.3	120	9	US-09-775-932-2
32	127.5	19.3	120	16	US-10-695-194-2
33	127.5	19.3	140	8	US-08-849-303-18
34	127.5	19.3	140	16	US-10-655-136-18
35	127.5	19.3	146	8	US-08-849-303-17
36	127.5	19.3	146	9	US-09-940-497-3
37	127.5	19.3	146	9	US-09-969-834-3
38	127.5	19.3	146	14	US-10-329-428-3
39	127.5	19.3	146	14	US-10-376-564-47
40	127.5	19.3	146	16	US-10-655-136-17
41	127.5	19.3	146	16	US-10-695-194-1
42	127.5	19.3	249	16	US-10-257-384A-4
43	127.5	19.3	641	16	US-10-257-384A-2
44	125	18.9	121	9	US-09-775-932-8
45	125	18.9	141	8	US-08-849-303-24

ALIGNMENTS

RESULT 1

US-10-162-335-82
; Sequence 82, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumsartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalte, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malyankar, Uriel M.
; APPLICANT: Miller, Isabelle
; APPLICANT: Padigaru, Muraidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shimkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Zeeb, Edward Z.
; APPLICANT: Voerhusen, Bryan D.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04

```
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 82
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-82

Query Match      100.0%; Score 660; DB 15; Length 390;
Best Local Similarity 100.0%; Pred. No. 2.9e-64;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 GKDVPPTKICVGCPRDPTNSPELEETLTHITTKNAENNATYFKIDNVKARQVW 60
Db      216 GKDVPPTKICVGCPRDPTNSPELEETLTHITTKNAENNATYFKIDNVKARQVW 275

Qy      61 AGKYFIDFVARETTCKESNEELTSCETKKGSLDCNAEVYVPWEKKIYPTVNCOP 120
Db      276 AGKYFIDFVARETTCKESNEELTSCETKKGSLDCNAEVYVPWEKKIYPTVNCOP 335

Qy      121 LGM 123
Db      336 LGM 338

RESULT 2
US-10-162-335-70
; Sequence 70, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalt, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malyankar, Uriel M.
; APPLICANT: Millet, Isabelle
; APPLICANT: Padigaru, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shinkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; APPLICANT: Zerhusen, Bryan D.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
```

```
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 70
; LENGTH: 398
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-70

Query Match      100.0%; Score 660; DB 15; Length 398;
Best Local Similarity 100.0%; Pred. No. 3e-64;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 GKDVPPTKICVGCPRDPTNSPELEETLTHITTKNAENNATYFKIDNVKARQVW 60
Db      224 GKDVPPTKICVGCPRDPTNSPELEETLTHITTKNAENNATYFKIDNVKARQVW 283

Qy      61 AGKYFIDFVARETTCKESNEELTSCETKKGSLDCNAEVYVPWEKKIYPTVNCOP 120
Db      284 AGKYFIDFVARETTCKESNEELTSCETKKGSLDCNAEVYVPWEKKIYPTVNCOP 343

Qy      121 LGM 123
Db      344 LGM 346

RESULT 3
US-09-919-039-29
; Sequence 29, Application US/09919039
; Publication No. US20030108871A1
; GENERAL INFORMATION:
; APPLICANT: Kaber, Matthew R.
; TITLE OF INVENTION: GENES EXPRESSED IN TREATED HUMAN C3A LIVER CELL CULTURES
; FILE REFERENCE: PA-0035 US
; CURRENT APPLICATION NUMBER: US/09/919,039
; CURRENT FILING DATE: 2002-09-09
; PRIOR APPLICATION NUMBER: 60/222,113
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 401
; SOFTWARE: PERL Program
; SEQ ID NO 29
; LENGTH: 427
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. US20030108871A1 167507CD1
US-09-919-039-29

Query Match      100.0%; Score 660; DB 10; Length 427;
Best Local Similarity 100.0%; Pred. No. 3.3e-64;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNATFYFKIDNVKARQVW 60
Db 253 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNATFYFKIDNVKARQVW 312
QY 61 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVYVPWEKKIYPTVNCOP 120
Db 333 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVYVPWEKKIYPTVNCOP 372
QY 121 LGM 123
Db 373 LGM 375

RESULT 4
US-10-162-335-72
; Sequence 72, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalt, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malvankar, Uriel M.
; APPLICANT: Milliet, Isabelle
; APPLICANT: Padigar, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shinkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 72
; LENGTH: 615
; TYPE: PRT
```

```
; ORGANISM: Homo sapiens
US-10-162-335-72
Query Match 100.0%; Score 660; DB 15; Length 615;
Best Local Similarity 100.0%; Pred. No. 5.4e-64; Indels 0; Gaps 0;
Matches 123; Conservative 0; Mismatches 0;
QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNATFYFKIDNVKARQVW 60
Db 224 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNATFYFKIDNVKARQVW 283
QY 61 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVYVPWEKKIYPTVNCOP 120
Db 284 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVYVPWEKKIYPTVNCOP 343
QY 121 LGM 123
Db 344 LGM 346

RESULT 5
US-10-162-335-74
; Sequence 74, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalt, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malvankar, Uriel M.
; APPLICANT: Milliet, Isabelle
; APPLICANT: Padigar, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shinkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
```

; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 74
; LENGTH: 644
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-74

Query Match 100.0%; Score 660; DB 15; Length 644;
Best Local Similarity 100.0%; Pred. No. 5.7e-64;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GKDFVQPPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQVQV 60
Db 253 GKDFVQPPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQVQV 312
Qy 61 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTVNCOP 120
Db 313 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTVNCOP 372
Qy 121 LGM 123
Db 373 LGM 375

RESULT 6

US-10-162-335-84

; Sequence 84, Application US/10162335

; Publication No. US20040009480A1

; GENERAL INFORMATION:

; APPLICANT: Anderson, David W.

; APPLICANT: Baumgartner, Jason C.

; APPLICANT: Boldog, Ferenc L.

; APPLICANT: Casman, Stacie J.

; APPLICANT: Edinger, Shalomit R.

; APPLICANT: Gangolli, Esha A.

; APPLICANT: Gerlach, Valerie

; APPLICANT: Gorman, Linda

; APPLICANT: Guo, Xiaojia (Sasha)

; APPLICANT: Hjalt, Tord

; APPLICANT: Kekuda, Ramesh

; APPLICANT: Li, Li

; APPLICANT: MacDougall, John R.

; APPLICANT: Malvankar, Uriel M.

; APPLICANT: Millet, Isabelle

; APPLICANT: Padigaru, Muralidhara

; APPLICANT: Patturajan, Meera

; APPLICANT: Pena, Carol E. A.

; APPLICANT: Rastelli, Luca

; APPLICANT: Shimkets, Richard A.

; APPLICANT: Stone, David J.

; APPLICANT: Spytek, Kimberly A.

; APPLICANT: Vernet, Corine A. M.

; APPLICANT: Voss, Edward Z.

; APPLICANT: Zerhusen, Bryan D.

; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method

; FILE REFERENCE: 21402-377 B

; CURRENT APPLICATION NUMBER: US/10/162,335

; CURRENT FILING DATE: 2002-10-01

; PRIOR APPLICATION NUMBER: 60/295,607

; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/295,661

; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/296,404

; PRIOR FILING DATE: 2001-06-06

; PRIOR APPLICATION NUMBER: 60/296,418

; PRIOR FILING DATE: 2001-06-06

; PRIOR APPLICATION NUMBER: 60/297,414

; PRIOR FILING DATE: 2001-06-11

; PRIOR APPLICATION NUMBER: 60/297,567

; PRIOR FILING DATE: 2001-06-12

; PRIOR APPLICATION NUMBER: 60/298,285

; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 84
; LENGTH: 644
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-84

Query Match 100.0%; Score 660; DB 15; Length 644;
Best Local Similarity 100.0%; Pred. No. 5.7e-64;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GKDFVQPPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQVQV 60
Db 253 GKDFVQPPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQVQV 312
Qy 61 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTVNCOP 120
Db 313 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTVNCOP 372
Qy 121 LGM 123
Db 373 LGM 375

RESULT 7

US-10-316-253-217

; Sequence 217, Application US/10316253

; Publication No. US20030162706A1

; GENERAL INFORMATION:

; APPLICANT: The Procter & Gamble Company

; APPLICANT: Peters, Kevin

; APPLICANT: Thompson, Larry

; APPLICANT: Wang, Feng

; APPLICANT: Greis, Kenneth

; TITLE OF INVENTION: Angiogenesis Modulating Proteins

; FILE REFERENCE: 8865M

; CURRENT APPLICATION NUMBER: US/10/316,253

; CURRENT FILING DATE: 2002-12-10

; PRIOR APPLICATION NUMBER: US 60/355,295

; PRIOR FILING DATE: 2002-02-08

; NUMBER OF SEQ ID NOS: 308

; SOFTWARE: Patentin version 3.1

; SEQ ID NO 217

; LENGTH: 424

; TYPE: PRT

; ORGANISM: Rattus norvegicus

US-10-316-253-217

Query Match 60.9%; Score 402; DB 14; Length 424;
Best Local Similarity 61.8%; Pred. No. 1.1e-35;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;

Qy 1 GKDFVQPPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQVQV 60
Db 246 GDDLPELLPNKCGCPREIPVDSPELKEALGHSLARLNAGHNIHFYFKIDTVKKAQVQV 305
Qy 61 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTVNCOP 120
Db 306 AGVIYVIEFIARETNCQSKQKTELTADCEKHLGSLNCNANVYMFENKVENKVVPTVRCOA 365
Qy 121 LGM 123
Db 366 LDM 368

```
RESULT 8
US-10-316-253-215
; Sequence 215, Application US/10316253
; Publication No. US20030162706A1
; GENERAL INFORMATION:
; APPLICANT: The Procter & Gamble Company
; APPLICANT: Peters, Kevin
; APPLICANT: Thompson, Larry
; APPLICANT: Wang, Feng
; APPLICANT: Greis, Kenneth
; TITLE OF INVENTION: Angiogenesis Modulating Proteins
; FILE REFERENCE: 8665M
; CURRENT APPLICATION NUMBER: US/10/316,253
; CURRENT FILING DATE: 2002-12-10
; PRIOR APPLICATION NUMBER: US 60/355,295
; PRIOR FILING DATE: 2002-02-08
; NUMBER OF SEQ ID NOS: 308
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 215
; LENGTH: 430
; TYPE: PRT
; ORGANISM: Rattus norvegicus
US-10-316-253-215

Query Match 60.8%; Score 401; DB 14; Length 430;
Best Local Similarity 61.8%; Pred. No. 1.5e-35;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;

QY 1 GKDFQPTKICVGCPRDIPNSPELEETLTHITKLAENNAATFYFKIDNVKARVQVW 60
Db 252 GDDLPELLPKRCGCPRIQVDSPELKEALGHSIAQLNAQHNHIFYFKIDTVKKAISQW 311

QY 61 AGKYFIDFVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVN 120
Db 312 AGVIIVIEFIARETNSKQSKTELTADCTKHLGSLNCNANVYMRPWENKVPVTRCQA 371

QY 121 LGM 123
Db 372 LDM 374

RESULT 9
US-09-969-834-1
; Sequence 1, Application US/09969834
; Patent No. US20020102711A1
; GENERAL INFORMATION:
; APPLICANT: Bandman, Olga
; TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESSES:
; ADDRESS: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/969,834
; FILING DATE: 01-Oct-2001
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/09/471,765
; FILING DATE: <Unknown>
; APPLICATION NUMBER: US/08/791,522
; FILING DATE: <Unknown>
; APPLICATION NUMBER: 09/471,765

FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0193 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 178 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
CLONE: 30443
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-969-834-1

Query Match 25.6%; Score 169; DB 9; Length 178;
Best Local Similarity 32.5%; Pred. No. 2.1e-10;
Matches 39; Conservative 22; Mismatches 49; Indels 10; Gaps 4;

QY 9 TKICVGCPRDIPNSPELEETLTHITKLAENNAATFYFKIDNVKARVQVWAGKYFID 68
Db 54 SRVKPGFPKTIKNDPGVLQAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKXMLE 113

QY 69 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVN--COPLM 123
Db 114 VEIGRTTCKNQHLRL-DDCDFQTNHTLQTLSCYSEVWVVPW---VPALRGACSPSL 168

RESULT 10
US-10-329-428-2
; Sequence 2, Application US/10329428
; Publication No. US20030114646A1
; GENERAL INFORMATION:
; APPLICANT: Li, et al.
; TITLE OF INVENTION: Human Cystatin F
; FILE REFERENCE: PF265PID2
; CURRENT APPLICATION NUMBER: US/10/329,428
; CURRENT FILING DATE: 2002-12-27
; PRIOR APPLICATION NUMBER: 60/014,795
; PRIOR FILING DATE: 1996-04-03
; PRIOR APPLICATION NUMBER: 08/832,535
; PRIOR FILING DATE: 1997-04-03
; PRIOR APPLICATION NUMBER: 09/019,485
; PRIOR FILING DATE: 1998-01-29
; PRIOR APPLICATION NUMBER: 09/528,436
; PRIOR FILING DATE: 2000-03-17
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 2
; LENGTH: 145
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-329-428-2

Query Match 25.1%; Score 165.5; DB 14; Length 145;
Best Local Similarity 32.5%; Pred. No. 3.9e-10;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;

QY 9 TKICVGCPRDIPNSPELEETLTHITKLAENNAATFYFKIDNVKARVQVWAGKYFID 68
Db 32 SRVKPGFPKTIKNDPGVLQAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKXMLE 91

QY 69 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKI-YPTVNC 118
Db 92 VEIGRTTCKNQHLRL-DDCDFQTNHTLQTLSCYSEVWVVPWLPLOHFEVPLRC 144

RESULT 11
US-09-746-783-197
```

; Sequence 197, Application US/09746783
; Publication No. US2003004935A1
; GENERAL INFORMATION:
; APPLICANT: Jacobs, Kenneth
; McCoy, John M.
; LaValle, Edward R.
; Racie, Lisa A.
; Treacy, Maurice
; Spaulding, Vikki
; Agostino, Michael J.
; Howes, Steven H.
; Fechtel, Kim
; TITLE OF INVENTION: SECRETED PROTEINS AND POLYNUCLEOTIDES
; ENCODING THEM
; NUMBER OF SEQUENCES: 231
; CORRESPONDENCE ADDRESS:
; ADDRESS: Genetics Institute, Inc.
; STREET: 87 CambridgePark Drive
; CITY: Cambridge
; STATE: MA
; COUNTRY: U.S.A.
; ZIP: 02140
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/746,783
; FILING DATE: 21-Dec-2000
; CLASSIFICATION: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Milasincic, Debra J.
; REGISTRATION NUMBER: 46,931
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 227-7400
; TELEFAX: (617) 742-4214
; INFORMATION FOR SEQ ID NO: 197:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 167 amino acids
; TYPE: amino acid
; STRANDEDNESS: <Unknown>
; TOPOLOGY: linear
; MOLECULE TYPE: Protein
; SEQUENCE DESCRIPTION: SEQ ID NO: 197:
US-09-746-783-197
Query Match 25.1%; Score 165.5; DB 10; Length 167;
Best Local Similarity 32.5%; Pred. No. 4.7e-10;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;
QY 9 TKICVCPDRDIPNPSPELEETLTHITKLNAENNAATFYFKIDNVKARVQVVGKYYFI 68
Db 54 SRVKGFPKTKTNDPGLQAARYSVKFNCTNDMLFKESRITRALVQIVKGLKYLE 113
QY 69 FVARETTCSKE---TKLGSGLDCAEVVVPWEKKI-YPTVNC 118
Db 114 VEIGRTCKKNQHLRL-DDCDQTNHTLAQTILSCSYSEVVVFWLQHFVEFVLR 166
RESULT 12
US-09-775-932-14
; Sequence 14, Application US/09775932
; Patent No. US20020137671A1
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503

; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 14
; LENGTH: 121
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-14
Query Match 21.0%; Score 138.5; DB 9; Length 121;
Best Local Similarity 31.5%; Pred. No. 3e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
QY 8 PTKICVCPDRDIPNPSPELEETLTHITKLNAENNAATFYFKIDNVKARVQVVGKYYFI 67
Db 2 PQERWVGLRDLSPDDPQVQKAAQAAVASYNMGNSIYYFRDTHIIKAOSQLVAGIKYFL 61
QY 68 DFVARETTCSKE---SNEELTESCETKKLGQ--SLDCNAEVVVPWE 109
Db 62 TMEVGTDCRKTGVTGDHVDLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 108
RESULT 13
US-09-775-932-12
; Sequence 12, Application US/09775932
; Patent No. US20020137671A1
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 12
; LENGTH: 128
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-12
Query Match 21.0%; Score 138.5; DB 9; Length 128;
Best Local Similarity 31.5%; Pred. No. 3.2e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
QY 8 PTKICVCPDRDIPNPSPELEETLTHITKLNAENNAATFYFKIDNVKARVQVVGKYYFI 67
Db 9 PQERWVGLRDLSPDDPQVQKAAQAAVASYNMGNSIYYFRDTHIIKAOSQLVAGIKYFL 68
QY 68 DFVARETTCSKE---SNEELTESCETKKLGQ--SLDCNAEVVVPWE 109
Db 69 TMEVGTDCRKTGVTGDHVDLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 115
RESULT 14
US-09-940-497-2
; Sequence 2, Application US/09940497
; Patent No. US20020052476A1
; GENERAL INFORMATION:
; APPLICANT: Ni et al.
; TITLE OF INVENTION: Human Cystatin E
; FILE REFERENCE: PE202P1D2
; CURRENT APPLICATION NUMBER: US/09/940,497
; CURRENT FILING DATE: 2001-08-29
; PRIOR APPLICATION NUMBER: US 09/241,376
; PRIOR FILING DATE: 1999-02-02
; PRIOR APPLICATION NUMBER: US 08/744,138
; PRIOR FILING DATE: 1996-11-05
; PRIOR APPLICATION NUMBER: US 08/461,030
; PRIOR FILING DATE: 1995-06-05

; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 2
; LENGTH: 149
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-940-497-2

Query Match 21.0%; Score 138.5; DB 9; Length 149;
Best Local Similarity 31.5%; Pred. No. 3.9e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 8 PTKICVCGPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARVQVWAGKKYFI 67

Db 30 PQRVVGRLDLPDPQVQKAAQAAVASYNMGNSIYFRDTHIIKQSLVAGIKYFL 89

QY 68 DFVARETTSKE-----SNEELTESCETKLGQ--SLDCNAEYVVPWE 109

Db 90 TMEGSTDCKRTRVTGHDVLT-TCPLAAGAQCEKRLCDFEVLVVPWQ 136

RESULT 15

US-08-849-303-16
; Sequence 16, Application US/08849303
; Publication No. US20030221209A1

GENERAL INFORMATION:

; APPLICANT: Atkinson, Howard J.
; APPLICANT: McPherson, Michael J.
; APPLICANT: Urwin, Peter E.

; TITLE OF INVENTION: MODIFIED PROTEINASE INHIBITORS

; NUMBER OF SEQUENCES: 79

; CORRESPONDENCE ADDRESS:

; ADDRESS: Klauber & Jackson

; STREET: 411 Hackensack Avenue, 4th Floor

; CITY: Hackensack

; STATE: New Jersey

; COUNTRY: USA

; ZIP: 07601

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/849.303

; FILING DATE: 21-MAY-1997

; CLASSIFICATION: 514

; ATTORNEY/AGENT INFORMATION:

; NAME: Jackson Esq., David A.

; REGISTRATION NUMBER: 26,742

; REFERENCE/DOCKET NUMBER: 1321-1-003

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: 201-487-5800

; TELEFAX: 201-343-1684

; TELEX: 133521

; INFORMATION FOR SEQ ID NO: 16:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 112 amino acids

; TYPE: amino acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: protein

; HYPOTHETICAL: NO

US-08-849-303-16

Query Match 20.6%; Score 136; DB 8; Length 112;
Best Local Similarity 30.6%; Pred. No. 5.1e-07;
Matches 34; Conservative 22; Mismatches 31; Indels 24; Gaps 5;

QY 22 NSPELEETLTHITKLAENNAATFYFKIDNVKKARVQVWAGKKYFIDFVARETTSKESN 81

Db 12 NEEGVQEALSPAVSBFNRKSRNDAYQSRVVRVRARQVVGMYFLDVELGRITCTK--S 69

QY 82 EELTESC-----ETKLGQSLDCNAEYVVPWEKKIYPTVN-----CQ 119
Db 70 QANLDSQCFHNPQHLKREKL-----CSFQYVYVPMWN-----TINLVKFSQ 111

Search completed: September 24, 2004, 14:13:03
Job time : 49.296 secs

